# CPC COOPERATIVE PATENT CLASSIFICATION

# C01B NON-METALLIC ELEMENTS; COMPOUNDS THEREOF; {METALLOIDS OR COMPOUNDS THEREOF NOT COVERED BY SUBCLASS C01C}

#### **NOTES**

- 1. In this subclass, tradenames that are often found in scientific and patent literature have been used in order to define precisely the scope of the groups.
- 2. Attention is drawn to the definitions of groups of chemical elements following the title of section C.

#### WARNINGS

1. The following IPC groups are not used in the CPC system. Subject matter covered by these groups is classified in the following CPC groups:

C01B 31/16 covered by <u>B01J 39/24, B01J 41/18</u> C01B 35/16, C01B 35/18 covered by <u>C01B 35/00</u> + s.gr.

2. General concordance IPC4 or IPC5 to IPC6 groups is as follows:

<u>C01B 25/37</u> (partly) : <u>C01B 37/002</u>

<u>C01B 25/453</u> : <u>C01B 37/00</u>, <u>C01B 39/54</u>

<u>C01B 33/185</u> : <u>C01B 37/02</u>

<u>C01B 33/28</u> and subgroups : <u>C01B 37/00</u>, <u>C01B 39/00</u>

<u>C01B 35/1009</u> : <u>C01B 37/06</u>, <u>C01B 39/12</u>, <u>C01B 39/54</u>

#### Hydrogen; Hydrides; Water; Synthesis gas from hydrocarbons

3/00 Hydrogen; Gaseous mixtures containing hydrogen; Separation of hydrogen from mixtures containing it (separation of gases by physical means B01D); Purification of hydrogen (production of water gas or synthesis gas from solid carbonaceous material C10J; purifying or modifying the chemical compositions of combustible technical gases containing carbon monoxide C10K)

## **NOTES**

- 1. In this group it is desirable to add the indexing codes of groups B01J 2208/00 and B01J 2219/00, for details relating to the reactors used in the generation of hydrogen or synthesis gas.
- 2. In groups C01B 3/12 C01B 3/18 and in groups C01B 3/22 C01B 3/586 it is desirable to add the indexing codes of group C01B 2203/00, for aspects relating to hydrogen or synthesis gas generation processes.
- 3/0005 {Reversible uptake of hydrogen by an appropriate medium, i.e. based on physical or chemical sorption phenomena or on reversible chemical reactions, e.g. for hydrogen storage purposes (purification of hydrogen C01B 3/508); Reversible gettering of hydrogen; Reversible uptake of hydrogen by electrodes}
- 3/001 Characterised by the uptaking medium; Treatment thereof}
- 3/0015 . . . {Organic compounds; Solutions thereof}

3/0021 . . . {Carbon, e.g. active carbon, carbon nanotubes, fullerenes; Treatment thereof}

3/0026 • • • {of one single metal or a rare earth metal; Treatment thereof}

**NOTES** 

- In all of the groups
   <u>C01B 3/0026</u> <u>C01B 3/0084</u>, the metallic
   storage materials may contain minor
   quantities of non-metals such as B, C,
   O, S, Se, Si; e.g. <u>C01B 3/0036</u> "only
   containing iron and titanium" includes Fe-Ti
   compositions comprising non-metals
- 2. In the groups C01B 3/0026 and C01B 3/0047 C01B 3/0068 a "rareearth metal" means one single metal or a combination of metals selected from the lanthanides, Sc or Y
- 3/0031 • {Intermetallic compounds; Metal alloys; Treatment thereof}
- 3/0036 . . . . {only containing iron and titanium; Treatment thereof}
- 3/0042 . . . . {only containing magnesium and nickel; Treatment thereof}
- 3/0047 . . . . {containing a rare earth metal; Treatment thereof}
- 3/0052 . . . . {also containing titanium} 3/0057 . . . . {also containing nickel}
- 3/0063 . . . . . {only containing a rare earth metal and only one other metal}
- 3/0068 . . . . . . {the other metal being nickel}
- 3/0073 . . . {Slurries, Suspensions}
- 3/0078 . . . {Composite solid storage mediums, i.e. coherent or loose mixtures of different solid constituents, chemically or structurally heterogeneous solid masses, coated solids or solids having a chemically modified surface region}
- 3/0084 . . . {Solid storage mediums characterised by their shape, e.g. pellets, sintered shaped bodies, sheets, porous compacts, spongy metals, hollow particles, solids with cavities, layered solids}

2/0000	(0.4	2/204	
3/0089	• {Ortho-para conversion}	3/384	• • • • {the catalyst being continuously externally
3/0094	• {Atomic hydrogen}	2/206	heated}
3/02	• Production of hydrogen or of gaseous mixtures	3/386	{Catalytic partial combustion}
2/025	containing {a substantial proportion of} hydrogen	3/388	• • • • {the heat being generated by superheated steam}
3/025	<ul> <li>{Preparation or purification of gas mixtures for ammonia synthesis}</li> </ul>	3/40	• • • • characterised by the catalyst
3/04	<ul> <li>by decomposition of inorganic compounds, e.g.</li> </ul>	3/40	using moving solid particles
3/04	ammonia {(C01B 3/0005 takes precedence)}	3/42	using moving solid particles using the fluidised bed technique
3/042	{Decomposition of water}	3/44	using the nutrised bed technique using discontinuously preheated non-moving
3/045	{in gaseous phase}	3/40	solid materials, e.g. blast and run
3/047	{Decomposition of ammonia}	3/48	• • • followed by reaction of water vapour with
3/06	by reaction of inorganic compounds containing	3/40	carbon monoxide
3/00	electro-positively bound hydrogen, e.g. water,	3/50	Separation of hydrogen or hydrogen containing
	acids, bases, ammonia, with inorganic reducing		gases from gaseous mixtures, e.g. purification
	agents (by electrolysis of water C25B 1/04)		(C01B 3/14 takes precedence)
3/061	• • • {by reaction of metal oxides with water}	3/501	• • {by diffusion}
3/063	{Cyclic methods}	3/503	• • {characterised by the membrane}
3/065	• • • {from a hydride}	3/505	{Membranes containing palladium}
3/066	• • • {by reaction of water with phosphorus}	3/506	• • {at low temperatures}
3/068	• • • {the hydrogen being generated from the water	3/508	• • {by selective and reversible uptake by an
	as a result of a cyclus of reactions, not covered		appropriate medium, i.e. the uptake being based
	by groups <u>C01B 3/063</u> or <u>C01B 3/105</u> }		on physical or chemical sorption phenomena or
3/08	with metals		on reversible chemical reactions (the appropriate
3/10	• • • by reaction of water vapour with metals		mediums <u>per se</u> <u>C01B 3/0005</u> )}
3/105	{Cyclic methods}	3/52	by contacting with liquids; Regeneration of used
3/12	• • • by reaction of water vapour with carbon	2/24	liquids {(C01B 3/508 takes precedence)}
	monoxide	3/54	including a catalytic reaction
3/14	Handling of heat and steam	3/56	• by contacting with solids; Regeneration of used
3/16	using catalysts	2/50	solids {(C01B 3/508 takes precedence)}
3/18	• • • using moving solid particles	3/58	including a catalytic reaction
3/20	• • by reaction of metal hydroxides with carbon	3/583	• • • • {the reaction being the selective oxidation of
	monoxide	2/586	carbon monoxide }
3/22	by decomposition of gaseous or liquid	3/586	carbon monoxide } {the reaction being a methanation reaction}
3/22	<ul> <li>by decomposition of gaseous or liquid organic compounds ({C01B 3/0005 takes</li> </ul>	3/586 <b>4/00</b>	{the reaction being a methanation reaction}  Hydrogen isotopes; Inorganic compounds thereof
3/22	<ul> <li>by decomposition of gaseous or liquid organic compounds ({C01B 3/0005 takes precedence }; coking liquid carbonaceous</li> </ul>		<ul> <li> {the reaction being a methanation reaction}</li> <li>Hydrogen isotopes; Inorganic compounds thereof prepared by isotope exchange, e.g. NH<sub>3</sub> + D<sub>2</sub></li> </ul>
	<ul> <li>by decomposition of gaseous or liquid organic compounds ({C01B 3/0005 takes precedence }; coking liquid carbonaceous materials C10B 55/00)</li> </ul>		<ul> <li> {the reaction being a methanation reaction}</li> <li>Hydrogen isotopes; Inorganic compounds thereof prepared by isotope exchange, e.g. NH<sub>3</sub> + D<sub>2</sub></li> <li>&gt; NH<sub>2</sub>D + HD (separation of isotopes B01D 59/00;</li> </ul>
3/24	<ul> <li>by decomposition of gaseous or liquid organic compounds ({C01B 3/0005 takes precedence } ; coking liquid carbonaceous materials C10B 55/00)</li> <li>of hydrocarbons</li> </ul>		Hydrogen isotopes; Inorganic compounds thereof prepared by isotope exchange, e.g. NH <sub>3</sub> + D <sub>2</sub> > NH <sub>2</sub> D + HD (separation of isotopes B01D 59/00; other chemical reactions to form compounds of
3/24 3/26	<ul> <li>by decomposition of gaseous or liquid organic compounds ({C01B 3/0005 takes precedence } ; coking liquid carbonaceous materials C10B 55/00)</li> <li>of hydrocarbons</li> <li>using catalysts</li> </ul>		Hydrogen isotopes; Inorganic compounds thereof prepared by isotope exchange, e.g. NH <sub>3</sub> + D <sub>2</sub> > NH <sub>2</sub> D + HD (separation of isotopes B01D 59/00; other chemical reactions to form compounds of hydrogen isotopes, see the relevant groups for
3/24 3/26 3/28	<ul> <li>by decomposition of gaseous or liquid organic compounds ({C01B 3/0005 takes precedence } ; coking liquid carbonaceous materials C10B 55/00)</li> <li>of hydrocarbons</li> <li>using catalysts</li> <li>using moving solid particles</li> </ul>		Hydrogen isotopes; Inorganic compounds thereof prepared by isotope exchange, e.g. NH <sub>3</sub> + D <sub>2</sub> > NH <sub>2</sub> D + HD (separation of isotopes B01D 59/00; other chemical reactions to form compounds of
3/24 3/26 3/28 3/30	<ul> <li>by decomposition of gaseous or liquid organic compounds ({C01B 3/0005 takes precedence } ; coking liquid carbonaceous materials C10B 55/00)</li> <li>of hydrocarbons</li> <li>using catalysts</li> <li>using moving solid particles</li> <li>using the fluidised bed technique</li> </ul>		Hydrogen isotopes; Inorganic compounds thereof prepared by isotope exchange, e.g. NH <sub>3</sub> + D <sub>2</sub> > NH <sub>2</sub> D + HD (separation of isotopes <u>B01D 59/00</u> ; other chemical reactions to form compounds of hydrogen isotopes, see the relevant groups for
3/24 3/26 3/28	<ul> <li>by decomposition of gaseous or liquid organic compounds ({C01B 3/0005 takes precedence } ; coking liquid carbonaceous materials C10B 55/00)</li> <li>of hydrocarbons</li> <li>using catalysts</li> <li>using moving solid particles</li> <li>using the fluidised bed technique</li> <li>by reaction of gaseous or liquid organic</li> </ul>	4/00	Hydrogen isotopes; Inorganic compounds thereof prepared by isotope exchange, e.g. NH <sub>3</sub> + D <sub>2</sub> > NH <sub>2</sub> D + HD (separation of isotopes B01D 59/00; other chemical reactions to form compounds of hydrogen isotopes, see the relevant groups for hydrogen compounds in class C01)
3/24 3/26 3/28 3/30	<ul> <li>by decomposition of gaseous or liquid organic compounds ({C01B 3/0005 takes precedence } ; coking liquid carbonaceous materials C10B 55/00)</li> <li>of hydrocarbons</li> <li>using catalysts</li> <li>using moving solid particles</li> <li>using the fluidised bed technique</li> </ul>	4/00 5/00	Hydrogen isotopes; Inorganic compounds thereof prepared by isotope exchange, e.g. NH <sub>3</sub> + D <sub>2</sub> > NH <sub>2</sub> D + HD (separation of isotopes B01D 59/00; other chemical reactions to form compounds of hydrogen isotopes, see the relevant groups for hydrogen compounds in class C01)  Water
3/24 3/26 3/28 3/30	<ul> <li>by decomposition of gaseous or liquid organic compounds ({C01B 3/0005 takes precedence } ; coking liquid carbonaceous materials C10B 55/00)</li> <li>of hydrocarbons</li> <li>using catalysts</li> <li>using moving solid particles</li> <li>using the fluidised bed technique</li> <li>by reaction of gaseous or liquid organic compounds with gasifying agents, e.g. water,</li> </ul>	4/00 5/00	Hydrogen isotopes; Inorganic compounds thereof prepared by isotope exchange, e.g. NH <sub>3</sub> + D <sub>2</sub> > NH <sub>2</sub> D + HD (separation of isotopes B01D 59/00; other chemical reactions to form compounds of hydrogen isotopes, see the relevant groups for hydrogen compounds in class C01)  Water  Heavy water; Preparation by chemical reaction of
3/24 3/26 3/28 3/30 3/32	<ul> <li>by decomposition of gaseous or liquid organic compounds ({C01B 3/0005 takes precedence } ; coking liquid carbonaceous materials C10B 55/00)</li> <li>of hydrocarbons</li> <li>using catalysts</li> <li>using moving solid particles</li> <li>using the fluidised bed technique</li> <li>by reaction of gaseous or liquid organic compounds with gasifying agents, e.g. water, carbon dioxide, air</li> </ul>	<b>4/00 5/00 5/02</b>	Hydrogen isotopes; Inorganic compounds thereof prepared by isotope exchange, e.g. NH <sub>3</sub> + D <sub>2</sub> > NH <sub>2</sub> D + HD (separation of isotopes B01D 59/00; other chemical reactions to form compounds of hydrogen isotopes, see the relevant groups for hydrogen compounds in class C01)  Water  Heavy water; Preparation by chemical reaction of hydrogen isotopes or their compounds, e.g. 4ND <sub>3</sub> + 7O <sub>2</sub> > 4NO <sub>2</sub> + 6D <sub>2</sub> O, 2D <sub>2</sub> + O <sub>2</sub> > 2D <sub>2</sub> O
3/24 3/26 3/28 3/30 3/32	<ul> <li>by decomposition of gaseous or liquid organic compounds ({C01B 3/0005 takes precedence}; coking liquid carbonaceous materials C10B 55/00)</li> <li>of hydrocarbons</li> <li>using catalysts</li> <li>using moving solid particles</li> <li>using the fluidised bed technique</li> <li>by reaction of gaseous or liquid organic compounds with gasifying agents, e.g. water, carbon dioxide, air</li> <li>{Catalytic reaction of gaseous or liquid organic</li> </ul>	4/00 5/00	<ul> <li>Hydrogen isotopes; Inorganic compounds thereof prepared by isotope exchange, e.g. NH<sub>3</sub> + D<sub>2</sub> &gt; NH<sub>2</sub>D + HD (separation of isotopes B01D 59/00; other chemical reactions to form compounds of hydrogen isotopes, see the relevant groups for hydrogen compounds in class C01)</li> <li>Water</li> <li>Heavy water; Preparation by chemical reaction of hydrogen isotopes or their compounds, e.g. 4ND<sub>3</sub> + 7O<sub>2</sub>&gt; 4NO<sub>2</sub> + 6D<sub>2</sub>O, 2D<sub>2</sub> + O<sub>2</sub>&gt; 2D<sub>2</sub>O</li> <li>Hydrides of metals (including fully or partially</li> </ul>
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3/24 3/26 3/28 3/30 3/32 3/323	<ul> <li>by decomposition of gaseous or liquid organic compounds ({CO1B 3/0005 takes precedence } ; coking liquid carbonaceous materials C10B 55/00)</li> <li>of hydrocarbons</li> <li>using catalysts</li> <li>using moving solid particles</li> <li>using the fluidised bed technique</li> <li>by reaction of gaseous or liquid organic compounds with gasifying agents, e.g. water, carbon dioxide, air</li> <li>{Catalytic reaction of gaseous or liquid organic compounds other than hydrocarbons with gasifying agents}</li> <li>{characterised by the catalyst}</li> <li>by reaction of hydrocarbons with gasifying agents</li> </ul>	<b>4/00 5/00 5/02</b>	<ul> <li>Hydrogen isotopes; Inorganic compounds thereof prepared by isotope exchange, e.g. NH<sub>3</sub> + D<sub>2</sub> &gt; NH<sub>2</sub>D + HD (separation of isotopes B01D 59/00; other chemical reactions to form compounds of hydrogen isotopes, see the relevant groups for hydrogen compounds in class C01)</li> <li>Water</li> <li>Heavy water; Preparation by chemical reaction of hydrogen isotopes or their compounds, e.g. 4ND<sub>3</sub> + 7O<sub>2</sub>&gt; 4NO<sub>2</sub> + 6D<sub>2</sub>O, 2D<sub>2</sub> + O<sub>2</sub>&gt; 2D<sub>2</sub>O</li> <li>Hydrides of metals (including fully or partially hydrided metals, alloys or intermetallic</li> </ul>
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3/24 3/26 3/28 3/30 3/32 3/323	<ul> <li>by decomposition of gaseous or liquid organic compounds ({CO1B 3/0005 takes precedence } ; coking liquid carbonaceous materials C10B 55/00)</li> <li>of hydrocarbons</li> <li>using catalysts</li> <li>using moving solid particles</li> <li>using the fluidised bed technique</li> <li>by reaction of gaseous or liquid organic compounds with gasifying agents, e.g. water, carbon dioxide, air</li> <li>{Catalytic reaction of gaseous or liquid organic compounds other than hydrocarbons with gasifying agents}</li> <li>{characterised by the catalyst}</li> <li>by reaction of hydrocarbons with gasifying agents</li> <li>with the aid of electrical means, electromagnetic or mechanical vibrations, or</li> </ul>	<b>4/00 5/00 5/02</b>	<ul> <li>Hydrogen isotopes; Inorganic compounds thereof prepared by isotope exchange, e.g. NH<sub>3</sub> + D<sub>2</sub> &gt; NH<sub>2</sub>D + HD (separation of isotopes B01D 59/00; other chemical reactions to form compounds of hydrogen isotopes, see the relevant groups for hydrogen compounds in class C01)</li> <li>Water         <ul> <li>Heavy water; Preparation by chemical reaction of hydrogen isotopes or their compounds, e.g. 4ND<sub>3</sub> + 7O<sub>2</sub>&gt; 4NO<sub>2</sub> + 6D<sub>2</sub>O, 2D<sub>2</sub> + O<sub>2</sub>&gt; 2D<sub>2</sub>O</li> <li>Hydrides of metals {including fully or partially hydrided metals, alloys or intermetallic compounds (use of some thereof for reversible sorption of hydrogen C01B 3/0005, C01B 3/508); Compounds containing at least one metal-</li> </ul> </li> </ul>
3/24 3/26 3/28 3/30 3/32 3/323 3/323 3/324 3/342	<ul> <li>by decomposition of gaseous or liquid organic compounds ({CO1B 3/0005 takes precedence } ; coking liquid carbonaceous materials C10B 55/00)</li> <li>of hydrocarbons</li> <li>using catalysts</li> <li>using moving solid particles</li> <li>using the fluidised bed technique</li> <li>by reaction of gaseous or liquid organic compounds with gasifying agents, e.g. water, carbon dioxide, air</li> <li>{Catalytic reaction of gaseous or liquid organic compounds other than hydrocarbons with gasifying agents}</li> <li>{characterised by the catalyst}</li> <li>by reaction of hydrocarbons with gasifying agents</li> <li>with the aid of electrical means, electromagnetic or mechanical vibrations, or particle radiations}</li> </ul>	<b>4/00 5/00 5/02</b>	<ul> <li>Hydrogen isotopes; Inorganic compounds thereof prepared by isotope exchange, e.g. NH<sub>3</sub> + D<sub>2</sub>&gt; NH<sub>2</sub>D + HD (separation of isotopes B01D 59/00; other chemical reactions to form compounds of hydrogen isotopes, see the relevant groups for hydrogen compounds in class C01)</li> <li>Water</li> <li>Heavy water; Preparation by chemical reaction of hydrogen isotopes or their compounds, e.g. 4ND<sub>3</sub> + 7O<sub>2</sub>&gt; 4NO<sub>2</sub> + 6D<sub>2</sub>O, 2D<sub>2</sub> + O<sub>2</sub>&gt; 2D<sub>2</sub>O</li> <li>Hydrides of metals {including fully or partially hydrided metals, alloys or intermetallic compounds (use of some thereof for reversible sorption of hydrogen C01B 3/0005, C01B 3/508); Compounds containing at least one metalhydrogen bond, e.g. (GeH<sub>3</sub>)<sub>2</sub>S, SiH GeH<sub>3</sub>;</li> <li>Monoborane or diborane; Addition complexes thereof (higher hydrides of boron, substituted</li> </ul>
3/24 3/26 3/28 3/30 3/32 3/323 3/323 3/324 3/344	<ul> <li>by decomposition of gaseous or liquid organic compounds ({C01B 3/0005 takes precedence } ; coking liquid carbonaceous materials C10B 55/00)</li> <li>of hydrocarbons</li> <li>using catalysts</li> <li>using moving solid particles</li> <li>using the fluidised bed technique</li> <li>by reaction of gaseous or liquid organic compounds with gasifying agents, e.g. water, carbon dioxide, air</li> <li>{Catalytic reaction of gaseous or liquid organic compounds other than hydrocarbons with gasifying agents}</li> <li>{characterised by the catalyst}</li> <li>by reaction of hydrocarbons with gasifying agents</li> <li>that catalytic reaction of particles with the aid of electrical means, electromagnetic or mechanical vibrations, or particle radiations}</li> <li>fusing non-catalytic solid particles}</li> </ul>	<b>5/00</b> 5/02 6/00	<ul> <li>Hydrogen isotopes; Inorganic compounds thereof prepared by isotope exchange, e.g. NH<sub>3</sub> + D<sub>2</sub> &gt; NH<sub>2</sub>D + HD (separation of isotopes B01D 59/00; other chemical reactions to form compounds of hydrogen isotopes, see the relevant groups for hydrogen compounds in class C01)</li> <li>Water         <ul> <li>Heavy water; Preparation by chemical reaction of hydrogen isotopes or their compounds, e.g. 4ND<sub>3</sub> + 7O<sub>2</sub>&gt; 4NO<sub>2</sub> + 6D<sub>2</sub>O, 2D<sub>2</sub> + O<sub>2</sub>&gt; 2D<sub>2</sub>O</li> <li>Hydrides of metals {including fully or partially hydrided metals, alloys or intermetallic compounds (use of some thereof for reversible sorption of hydrogen C01B 3/0005, C01B 3/508); Compounds containing at least one metalhydrogen bond, e.g. (GeH<sub>3</sub>)<sub>2</sub>S, SiH GeH};</li> <li>Monoborane or diborane; Addition complexes thereof (higher hydrides of boron, substituted hydrides of boron C01B 35/00)</li> </ul> </li> </ul>
3/24 3/26 3/28 3/30 3/32 3/323 3/326 3/34 3/342	<ul> <li>by decomposition of gaseous or liquid organic compounds ({C01B 3/0005 takes precedence } ; coking liquid carbonaceous materials C10B 55/00)</li> <li>of hydrocarbons</li> <li>using catalysts</li> <li>using moving solid particles</li> <li>using the fluidised bed technique</li> <li>by reaction of gaseous or liquid organic compounds with gasifying agents, e.g. water, carbon dioxide, air</li> <li>{Catalytic reaction of gaseous or liquid organic compounds other than hydrocarbons with gasifying agents}</li> <li>{characterised by the catalyst}</li> <li>by reaction of hydrocarbons with gasifying agents</li> <li>that catalyst is agent is agents</li> <li>that catalyst is agents</li> <li>that catalyst is agent is agent</li></ul>	<b>4/00 5/00 5/02</b>	Hydrogen isotopes; Inorganic compounds thereof prepared by isotope exchange, e.g. NH <sub>3</sub> + D <sub>2</sub> > NH <sub>2</sub> D + HD (separation of isotopes B01D 59/00; other chemical reactions to form compounds of hydrogen isotopes, see the relevant groups for hydrogen compounds in class C01)  Water  Heavy water; Preparation by chemical reaction of hydrogen isotopes or their compounds, e.g. 4ND <sub>3</sub> + 7O <sub>2</sub> > 4NO <sub>2</sub> + 6D <sub>2</sub> O, 2D <sub>2</sub> + O <sub>2</sub> > 2D <sub>2</sub> O  Hydrides of metals {including fully or partially hydrided metals, alloys or intermetallic compounds (use of some thereof for reversible sorption of hydrogen C01B 3/0005, C01B 3/508); Compounds containing at least one metalhydrogen bond, e.g. (GeH <sub>3</sub> ) <sub>2</sub> S, SiH GeH}; Monoborane or diborane; Addition complexes thereof (higher hydrides of boron, substituted hydrides of boron C01B 35/00)  Hydrides containing only one metal and one or
3/24 3/26 3/28 3/30 3/32 3/323 3/323 3/324 3/344	<ul> <li>by decomposition of gaseous or liquid organic compounds ({CO1B 3/0005 takes precedence } ; coking liquid carbonaceous materials C10B 55/00)</li> <li>of hydrocarbons</li> <li>using catalysts</li> <li>using moving solid particles</li> <li>using the fluidised bed technique</li> <li>by reaction of gaseous or liquid organic compounds with gasifying agents, e.g. water, carbon dioxide, air</li> <li>{Catalytic reaction of gaseous or liquid organic compounds other than hydrocarbons with gasifying agents}</li> <li>{characterised by the catalyst}</li> <li>by reaction of hydrocarbons with gasifying agents</li> <li>{with the aid of electrical means, electromagnetic or mechanical vibrations, or particle radiations}</li> <li>{using non-catalytic solid particles}</li> <li>{using heat generated by superheated steam}</li> <li>{by direct contact with heat accumulating</li> </ul>	<b>5/00</b> 5/02 <b>6/00</b>	<ul> <li>Hydrogen isotopes; Inorganic compounds thereof prepared by isotope exchange, e.g. NH<sub>3</sub> + D<sub>2</sub> &gt; NH<sub>2</sub>D + HD (separation of isotopes B01D 59/00; other chemical reactions to form compounds of hydrogen isotopes, see the relevant groups for hydrogen compounds in class C01)</li> <li>Water</li> <li>Heavy water; Preparation by chemical reaction of hydrogen isotopes or their compounds, e.g. 4ND<sub>3</sub> + 7O<sub>2</sub>&gt; 4NO<sub>2</sub> + 6D<sub>2</sub>O, 2D<sub>2</sub> + O<sub>2</sub>&gt; 2D<sub>2</sub>O</li> <li>Hydrides of metals {including fully or partially hydrided metals, alloys or intermetallic compounds (use of some thereof for reversible sorption of hydrogen C01B 3/0005, C01B 3/508); Compounds containing at least one metalhydrogen bond, e.g. (GeH<sub>3</sub>)<sub>2</sub>S, SiH GeH}; Monoborane or diborane; Addition complexes thereof (higher hydrides of boron, substituted hydrides containing only one metal and one or several non-metals}</li> </ul>
3/24 3/26 3/28 3/30 3/32 3/323 3/326 3/34 3/342 3/344 3/346 3/348	<ul> <li>by decomposition of gaseous or liquid organic compounds ({CO1B 3/0005 takes precedence } ; coking liquid carbonaceous materials C10B 55/00)</li> <li>of hydrocarbons</li> <li>using catalysts</li> <li>using moving solid particles</li> <li>using the fluidised bed technique</li> <li>by reaction of gaseous or liquid organic compounds with gasifying agents, e.g. water, carbon dioxide, air</li> <li>{Catalytic reaction of gaseous or liquid organic compounds other than hydrocarbons with gasifying agents}</li> <li>{characterised by the catalyst}</li> <li>by reaction of hydrocarbons with gasifying agents</li> <li>{with the aid of electrical means, electromagnetic or mechanical vibrations, or particle radiations}</li> <li>{using non-catalytic solid particles}</li> <li>{using heat generated by superheated steam}</li> <li>{by direct contact with heat accumulating liquids, e.g. molten metals, molten salts}</li> </ul>	<b>4/00 5/00 5/02 6/00</b> 6/003 6/006	<ul> <li>Hydrogen isotopes; Inorganic compounds thereof prepared by isotope exchange, e.g. NH<sub>3</sub> + D<sub>2</sub>&gt; NH<sub>2</sub>D + HD (separation of isotopes B01D 59/00; other chemical reactions to form compounds of hydrogen isotopes, see the relevant groups for hydrogen compounds in class C01)</li> <li>Water</li> <li>Heavy water; Preparation by chemical reaction of hydrogen isotopes or their compounds, e.g. 4ND<sub>3</sub> + 7O<sub>2</sub>&gt; 4NO<sub>2</sub> + 6D<sub>2</sub>O, 2D<sub>2</sub> + O<sub>2</sub>&gt; 2D<sub>2</sub>O</li> <li>Hydrides of metals {including fully or partially hydrided metals, alloys or intermetallic compounds (use of some thereof for reversible sorption of hydrogen C01B 3/0005, C01B 3/508); Compounds containing at least one metalhydrogen bond, e.g. (GeH<sub>3</sub>)<sub>2</sub>S, SiH GeH}; Monoborane or diborane; Addition complexes thereof (higher hydrides of boron, substituted hydrides of boron C01B 35/00)</li> <li>{Hydrides containing only one metal and one or several non-metals}</li> <li>{only one metal and one or several halogens}</li> </ul>
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3/24 3/26 3/28 3/30 3/32 3/323 3/323 3/326 3/344 3/346 3/348 3/36	<ul> <li>by decomposition of gaseous or liquid organic compounds ({CO1B 3/0005 takes precedence } ; coking liquid carbonaceous materials C10B 55/00)</li> <li>of hydrocarbons</li> <li>using catalysts</li> <li>using moving solid particles</li> <li>using the fluidised bed technique</li> <li>by reaction of gaseous or liquid organic compounds with gasifying agents, e.g. water, carbon dioxide, air</li> <li>{Catalytic reaction of gaseous or liquid organic compounds other than hydrocarbons with gasifying agents}</li> <li>{characterised by the catalyst}</li> <li>by reaction of hydrocarbons with gasifying agents</li> <li>{with the aid of electrical means, electromagnetic or mechanical vibrations, or particle radiations}</li> <li>{using non-catalytic solid particles}</li> <li>{using heat generated by superheated steam}</li> <li>{by direct contact with heat accumulating liquids, e.g. molten metals, molten salts}</li> <li>using oxygen or mixtures containing oxygen as gasifying agents</li> </ul>	<b>5/00</b> 5/02 <b>6/00</b> 6/003  6/006  6/02	<ul> <li>Hydrogen isotopes; Inorganic compounds thereof prepared by isotope exchange, e.g. NH<sub>3</sub> + D<sub>2</sub>&gt; NH<sub>2</sub>D + HD (separation of isotopes B01D 59/00; other chemical reactions to form compounds of hydrogen isotopes, see the relevant groups for hydrogen compounds in class C01)</li> <li>Water</li> <li>Heavy water; Preparation by chemical reaction of hydrogen isotopes or their compounds, e.g. 4ND<sub>3</sub> + 7O<sub>2</sub>&gt; 4NO<sub>2</sub> + 6D<sub>2</sub>O, 2D<sub>2</sub> + O<sub>2</sub>&gt; 2D<sub>2</sub>O</li> <li>Hydrides of metals {including fully or partially hydrided metals, alloys or intermetallic compounds (use of some thereof for reversible sorption of hydrogen C01B 3/0005, C01B 3/508); Compounds containing at least one metalhydrogen bond, e.g. (GeH<sub>3</sub>)<sub>2</sub>S, SiH GeH}; Monoborane or diborane; Addition complexes thereof (higher hydrides of boron, substituted hydrides of boron C01B 35/00)</li> <li>{Hydrides containing only one metal and one or several non-metals}</li> <li>• {only one metal and one or several halogens}</li> <li>• Hydrides of transition elements; Addition complexes thereof</li> </ul>
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6/06	TT 1:1 C 1 :: 11: : 1: 4 11:	7/07/12	( ( 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1
6/06	<ul> <li>Hydrides of aluminium, gallium, indium, thallium, germanium, tin, lead, arsenic, antimony, bismuth</li> </ul>	7/0743	• • { of gaseous or dissolved chlorine }
	or polonium; Monoborane; Diborane; Addition	7/075	of liquid chlorine
	complexes thereof	7/09	Bromine; Hydrogen bromide
6/065	• • {Hydrides of arsenic or antimony}	7/093 7/096	• • {Hydrogen bromide}
6/10	Monoborane; Diborane; Addition complexes		{Bromine}
0/10	thereof	7/13	• Iodine; Hydrogen iodide
6/11	Preparation from boron or inorganic	7/135	• • {Hydrogen iodide}
	compounds containing boron and oxygen	7/14	Iodine
6/13	Addition complexes of monoborane or	7/16	Preparation from seaweed
	diborane, e.g. with phosphine, arsine or	7/19	• Fluorine; Hydrogen fluoride
	hydrazine	7/191	• • {Hydrogen fluoride}
6/15	Metal borohydrides; Addition complexes	7/192	{Preparation from fluorspar}
	thereof	7/193	Preparation from silicon tetrafluoride, fluosilicic acid or fluosilicates}
6/17	Preparation from boron or inorganic	7/194	{Preparation from ammonium fluoride}
	compounds containing boron and oxygen	7/194	
6/19	Preparation from other compounds of	7/193 7/196	<ul><li> {Separation; Purification}</li><li> {by distillation}</li></ul>
	boron		The state of the s
6/21	Preparation of borohydrides of	7/197	• • • {by adsorption}
	alkali metals, alkaline earth metals,	7/198	• • • {by solid ion-exchangers}
	magnesium or beryllium; Addition	7/20	. Fluorine
	complexes thereof, e.g. LiBH <sub>4.2</sub> N <sub>2</sub> H <sub>4</sub> ,	7/24	Inter-halogen compounds
6/00	NaB <sub>2</sub> H <sub>7</sub>	9/00	General methods of preparing halides (particular
6/23	Preparation of borohydrides of other		individual halides, see the relevant groups in
	metals, e.g. aluminium borohydride;		<u>C01B</u> - <u>C01G</u> according to the element combined
	Addition complexes thereof, e.g. $Li[Al(BH_4)_3H]$		with the halogen; electrolytic production of inorganic
6/24	Hydrides containing at least two metals; Addition		compounds <u>C25B</u> )
0/24	complexes thereof (C01B 6/13 - C01B 6/23 take	9/02	• Chlorides
	precedence)	9/04	• Bromides
6/243	• • {containing only hydrogen, aluminium and alkali	9/06	• Iodides
0, 2 . 5	metals, e.g. Li(AlH <sub>4</sub> )}	9/08	• Fluorides
6/246	• {also containing non-metals other than hydrogen}	11/00	Ovides or avvacide of halogens: Salte thereof
6/246 6/26	<ul> <li>. {also containing non-metals other than hydrogen}</li> <li>. Preparation from the metal with the highest</li> </ul>	<b>11/00</b>	Oxides or oxyacids of halogens; Salts thereof
6/246 6/26	Preparation from the metal with the highest	11/02	• Oxides of chlorine
	Preparation from the metal with the highest valency or from its oxides or salts of its oxyacids	11/02 11/021	<ul> <li>Oxides of chlorine</li> <li>{Chlorine hemioxide (Cl<sub>2</sub>O)}</li> </ul>
6/26	Preparation from the metal with the highest	11/02 11/021 11/022	<ul> <li>Oxides of chlorine</li> <li>{Chlorine hemioxide (Cl<sub>2</sub>O)}</li> <li>{Chlorine dioxide (ClO<sub>2</sub>)}</li> </ul>
6/26 6/34	Preparation from the metal with the highest valency or from its oxides or salts of its oxyacids	11/02 11/021 11/022 11/023	<ul> <li>Oxides of chlorine</li> <li>{Chlorine hemioxide (Cl<sub>2</sub>O)}</li> <li>{Chlorine dioxide (ClO<sub>2</sub>)}</li> <li>{Preparation from chlorites or chlorates}</li> </ul>
6/26 6/34 <b>Halogens, co</b>	<ul> <li>Preparation from the metal with the highest valency or from its oxides or salts of its oxyacids</li> <li>Purification; Stabilisation</li> </ul>	11/02 11/021 11/022 11/023 11/024	<ul> <li>Oxides of chlorine</li> <li>{Chlorine hemioxide (Cl<sub>2</sub>O)}</li> <li>{Chlorine dioxide (ClO<sub>2</sub>)}</li> <li>{Preparation from chlorites or chlorates}</li> <li>{from chlorites}</li> </ul>
6/26 6/34 <b>Halogens, co</b> <b>7/00</b>	Preparation from the metal with the highest valency or from its oxides or salts of its oxyacids     Purification; Stabilisation  Ompounds thereof  Halogens; Halogen acids (oxyacids C01B 11/00)	11/02 11/021 11/022 11/023	<ul> <li>Oxides of chlorine</li> <li>{Chlorine hemioxide (Cl<sub>2</sub>O)}</li> <li>{Chlorine dioxide (ClO<sub>2</sub>)}</li> <li>{Preparation from chlorites or chlorates}</li> <li>{from chlorites}</li> <li>{from chlorates without any other reaction}</li> </ul>
6/26 6/34 Halogens, co 7/00 7/01	Preparation from the metal with the highest valency or from its oxides or salts of its oxyacids     Purification; Stabilisation  mpounds thereof  Halogens; Halogen acids (oxyacids C01B 11/00)     Chlorine; Hydrogen chloride	11/02 11/021 11/022 11/023 11/024 11/025	<ul> <li>Oxides of chlorine</li> <li>{Chlorine hemioxide (Cl<sub>2</sub>O)}</li> <li>{Chlorine dioxide (ClO<sub>2</sub>)}</li> <li>{Preparation from chlorites or chlorates}</li> <li>{from chlorites}</li> <li>{from chlorates without any other reaction reducing agent than chloride ions}</li> </ul>
6/26 6/34 <b>Halogens, co</b> <b>7/00</b>	Preparation from the metal with the highest valency or from its oxides or salts of its oxyacids     Purification; Stabilisation  mpounds thereof  Halogens; Halogen acids (oxyacids C01B 11/00)     Chlorine; Hydrogen chloride     Preparation of hydrogen chloride from the	11/02 11/021 11/022 11/023 11/024	<ul> <li>Oxides of chlorine</li> <li>{Chlorine hemioxide (Cl<sub>2</sub>O)}</li> <li>{Chlorine dioxide (ClO<sub>2</sub>)}</li> <li>{Preparation from chlorites or chlorates}</li> <li>{from chlorites}</li> <li>{from chlorates without any other reaction reducing agent than chloride ions}</li> <li>{from chlorate ions in the presence of a</li> </ul>
6/26 6/34 Halogens, co 7/00 7/01 7/012	Preparation from the metal with the highest valency or from its oxides or salts of its oxyacids     Purification; Stabilisation  mpounds thereof  Halogens; Halogen acids (oxyacids C01B 11/00)     Chlorine; Hydrogen chloride     Preparation of hydrogen chloride from the elements}	11/02 11/021 11/022 11/023 11/024 11/025	<ul> <li>Oxides of chlorine</li> <li>{Chlorine hemioxide (Cl<sub>2</sub>O)}</li> <li>{Chlorine dioxide (ClO<sub>2</sub>)}</li> <li>{Preparation from chlorites or chlorates}</li> <li>{from chlorites}</li> <li>{from chlorates without any other reaction reducing agent than chloride ions}</li> <li>{from chlorate ions in the presence of a peroxidic compound, e.g. hydrogen peroxide,</li> </ul>
6/26 6/34 Halogens, co 7/00 7/01	<ul> <li>Preparation from the metal with the highest valency or from its oxides or salts of its oxyacids</li> <li>Purification; Stabilisation</li> <li>Purification; Stabilisation</li> <li>Palogens; Halogen acids (oxyacids C01B 11/00)</li> <li>Chlorine; Hydrogen chloride</li> <li>{Preparation of hydrogen chloride from the elements}</li> <li>{Chlorine hydrates; Obtaining chlorine</li> </ul>	11/02 11/021 11/022 11/023 11/024 11/025	<ul> <li>Oxides of chlorine</li> <li>{Chlorine hemioxide (Cl<sub>2</sub>O)}</li> <li>{Chlorine dioxide (ClO<sub>2</sub>)}</li> <li>{Preparation from chlorites or chlorates}</li> <li>{from chlorites}</li> <li>{from chlorates without any other reaction reducing agent than chloride ions}</li> <li>{from chlorate ions in the presence of a peroxidic compound, e.g. hydrogen peroxide, ozone, peroxysulfates}</li> </ul>
6/26 6/34 Halogens, co 7/00 7/01 7/012 7/015	<ul> <li>Preparation from the metal with the highest valency or from its oxides or salts of its oxyacids</li> <li>Purification; Stabilisation</li> <li>Purification; Stabilisation</li> <li>Palogens; Halogen acids (oxyacids C01B 11/00)</li> <li>Chlorine; Hydrogen chloride</li> <li>{Preparation of hydrogen chloride from the elements}</li> <li>{Chlorine hydrates; Obtaining chlorine therefrom}</li> </ul>	11/02 11/021 11/022 11/023 11/024 11/025	<ul> <li>Oxides of chlorine</li> <li>{Chlorine hemioxide (Cl<sub>2</sub>O)}</li> <li>{Chlorine dioxide (ClO<sub>2</sub>)}</li> <li>{Preparation from chlorites or chlorates}</li> <li>{from chlorites}</li> <li>{from chlorates without any other reaction reducing agent than chloride ions}</li> <li>{from chlorate ions in the presence of a peroxidic compound, e.g. hydrogen peroxide, ozone, peroxysulfates}</li> <li>{from chlorate ions in the presence of</li> </ul>
6/26 6/34 Halogens, co 7/00 7/01 7/012	<ul> <li>Preparation from the metal with the highest valency or from its oxides or salts of its oxyacids</li> <li>Purification; Stabilisation</li> <li>Purification; Stabilisation</li> <li>Purification; Stabilisation</li> <li>Purification; Stabilisation</li> <li>Compounds thereof</li> <li>Halogens; Halogen acids (oxyacids C01B 11/00)</li> <li>Chlorine; Hydrogen chloride</li> <li>{Preparation of hydrogen chloride from the elements}</li> <li>{Chlorine hydrates; Obtaining chlorine therefrom}</li> <li>{Preparation of hydrogen chloride by reacting</li> </ul>	11/02 11/021 11/022 11/023 11/024 11/025	<ul> <li>Oxides of chlorine</li> <li>{Chlorine hemioxide (Cl<sub>2</sub>O)}</li> <li>{Chlorine dioxide (ClO<sub>2</sub>)}</li> <li>{Preparation from chlorites or chlorates}</li> <li>{from chlorites}</li> <li>{from chlorates without any other reaction reducing agent than chloride ions}</li> <li>{from chlorate ions in the presence of a peroxidic compound, e.g. hydrogen peroxide, ozone, peroxysulfates}</li> </ul>
6/26 6/34 Halogens, co 7/00 7/01 7/012 7/015	<ul> <li>Preparation from the metal with the highest valency or from its oxides or salts of its oxyacids</li> <li>Purification; Stabilisation</li> <li>Impounds thereof</li> <li>Halogens; Halogen acids (oxyacids C01B 11/00)</li> <li>Chlorine; Hydrogen chloride</li> <li>{Preparation of hydrogen chloride from the elements}</li> <li>{Chlorine hydrates; Obtaining chlorine therefrom}</li> <li>{Preparation of hydrogen chloride by reacting together chlorine, water and carbon or carbon</li> </ul>	11/02 11/021 11/022 11/023 11/024 11/025	<ul> <li>Oxides of chlorine</li> <li>{Chlorine hemioxide (Cl<sub>2</sub>O)}</li> <li>{Chlorine dioxide (ClO<sub>2</sub>)}</li> <li>{Preparation from chlorites or chlorates}</li> <li>{from chlorites}</li> <li>{from chlorates without any other reaction reducing agent than chloride ions}</li> <li>{from chlorate ions in the presence of a peroxidic compound, e.g. hydrogen peroxide, ozone, peroxysulfates}</li> <li>{from chlorate ions in the presence of a nitrogen compound selected from</li> </ul>
6/26 6/34 Halogens, co 7/00 7/01 7/012 7/015	<ul> <li>Preparation from the metal with the highest valency or from its oxides or salts of its oxyacids</li> <li>Purification; Stabilisation</li> <li>Purification; Stabilisation</li> <li>Purification; Stabilisation</li> <li>Purification; Stabilisation</li> <li>Compounds thereof</li> <li>Halogens; Halogen acids (oxyacids C01B 11/00)</li> <li>Chlorine; Hydrogen chloride</li> <li>{Preparation of hydrogen chloride from the elements}</li> <li>{Chlorine hydrates; Obtaining chlorine therefrom}</li> <li>{Preparation of hydrogen chloride by reacting</li> </ul>	11/02 11/021 11/022 11/023 11/024 11/025	<ul> <li>Oxides of chlorine</li> <li>{Chlorine hemioxide (Cl<sub>2</sub>O)}</li> <li>{Chlorine dioxide (ClO<sub>2</sub>)}</li> <li>{Preparation from chlorites or chlorates}</li> <li>{from chlorites}</li> <li>{from chlorates without any other reaction reducing agent than chloride ions}</li> <li>{from chlorate ions in the presence of a peroxidic compound, e.g. hydrogen peroxide, ozone, peroxysulfates}</li> <li>{from chlorate ions in the presence of a nitrogen compound selected from nitrogen dioxide, nitrate or nitrite ions,</li> </ul>
6/26 6/34 Halogens, co 7/00 7/01 7/012 7/015	<ul> <li>Preparation from the metal with the highest valency or from its oxides or salts of its oxyacids</li> <li>Purification; Stabilisation</li> <li>Impounds thereof</li> <li>Halogens; Halogen acids (oxyacids C01B 11/00)</li> <li>Chlorine; Hydrogen chloride</li> <li>{Preparation of hydrogen chloride from the elements}</li> <li>{Chlorine hydrates; Obtaining chlorine therefrom}</li> <li>{Preparation of hydrogen chloride by reacting together chlorine, water and carbon or carbon monoxide (the carbon not acting only as</li> </ul>	11/02 11/021 11/022 11/023 11/024 11/025	<ul> <li>Oxides of chlorine</li> <li>{Chlorine hemioxide (Cl<sub>2</sub>O)}</li> <li>{Chlorine dioxide (ClO<sub>2</sub>)}</li> <li>{Preparation from chlorites or chlorates}</li> <li>{from chlorites}</li> <li>{from chlorates without any other reaction reducing agent than chloride ions}</li> <li>{from chlorate ions in the presence of a peroxidic compound, e.g. hydrogen peroxide, ozone, peroxysulfates}</li> <li>{from chlorate ions in the presence of a nitrogen compound selected from nitrogen dioxide, nitrate or nitrite ions, nitrosylchloride, hydrazine or hydrazine</li> </ul>
6/26 6/34 Halogens, co 7/00 7/01 7/012 7/015 7/017	<ul> <li>Preparation from the metal with the highest valency or from its oxides or salts of its oxyacids</li> <li>Purification; Stabilisation</li> <li>Impounds thereof</li> <li>Halogens; Halogen acids (oxyacids C01B 11/00)</li> <li>Chlorine; Hydrogen chloride</li> <li>{Preparation of hydrogen chloride from the elements}</li> <li>{Chlorine hydrates; Obtaining chlorine therefrom}</li> <li>{Preparation of hydrogen chloride by reacting together chlorine, water and carbon or carbon monoxide (the carbon not acting only as catalyst)}</li> </ul>	11/02 11/021 11/022 11/023 11/024 11/025 11/026	<ul> <li>Oxides of chlorine</li> <li>{Chlorine hemioxide (Cl<sub>2</sub>O)}</li> <li>{Chlorine dioxide (ClO<sub>2</sub>)}</li> <li>{Preparation from chlorites or chlorates}</li> <li>{from chlorites}</li> <li>{from chlorates without any other reaction reducing agent than chloride ions}</li> <li>{from chlorate ions in the presence of a peroxidic compound, e.g. hydrogen peroxide, ozone, peroxysulfates}</li> <li>{from chlorate ions in the presence of a nitrogen compound selected from nitrogen dioxide, nitrate or nitrite ions, nitrosylchloride, hydrazine or hydrazine compounds}</li> </ul>
6/26 6/34 Halogens, co 7/00 7/01 7/012 7/015 7/017	<ul> <li>Preparation from the metal with the highest valency or from its oxides or salts of its oxyacids</li> <li>Purification; Stabilisation</li> <li>Impounds thereof</li> <li>Halogens; Halogen acids (oxyacids C01B 11/00)</li> <li>Chlorine; Hydrogen chloride</li> <li>{Preparation of hydrogen chloride from the elements}</li> <li>{Chlorine hydrates; Obtaining chlorine therefrom}</li> <li>{Preparation of hydrogen chloride by reacting together chlorine, water and carbon or carbon monoxide (the carbon not acting only as catalyst)}</li> <li>Preparation from chlorides</li> </ul>	11/02 11/021 11/022 11/023 11/024 11/025 11/026	<ul> <li>Oxides of chlorine</li> <li>{Chlorine hemioxide (Cl<sub>2</sub>O)}</li> <li>{Chlorine dioxide (ClO<sub>2</sub>)}</li> <li>{Preparation from chlorites or chlorates}</li> <li>{from chlorites}</li> <li>{from chlorates without any other reaction reducing agent than chloride ions}</li> <li>{from chlorate ions in the presence of a peroxidic compound, e.g. hydrogen peroxide, ozone, peroxysulfates}</li> <li>{from chlorate ions in the presence of a nitrogen compound selected from nitrogen dioxide, nitrate or nitrite ions, nitrosylchloride, hydrazine or hydrazine compounds}</li> <li>{Separation; Purification}</li> </ul>
6/26 6/34 Halogens, co 7/00 7/01 7/012 7/015 7/017	<ul> <li>Preparation from the metal with the highest valency or from its oxides or salts of its oxyacids</li> <li>Purification; Stabilisation</li> <li>Purification; Stabilisation</li> <li>Preparation; Halogen acids (oxyacids C01B 11/00)</li> <li>Chlorine; Hydrogen chloride</li> <li>{Preparation of hydrogen chloride from the elements}</li> <li>{Chlorine hydrates; Obtaining chlorine therefrom}</li> <li>{Preparation of hydrogen chloride by reacting together chlorine, water and carbon or carbon monoxide (the carbon not acting only as catalyst)}</li> <li>Preparation from chlorides</li> <li>{Preparation of hydrogen chloride from</li> </ul>	11/02 11/021 11/022 11/023 11/024 11/025 11/026	<ul> <li>Oxides of chlorine</li> <li>{Chlorine hemioxide (Cl<sub>2</sub>O)}</li> <li>{Chlorine dioxide (ClO<sub>2</sub>)}</li> <li>{Preparation from chlorites or chlorates}</li> <li>{from chlorites}</li> <li>{from chlorates without any other reaction reducing agent than chloride ions}</li> <li>{from chlorate ions in the presence of a peroxidic compound, e.g. hydrogen peroxide, ozone, peroxysulfates}</li> <li>{from chlorate ions in the presence of a nitrogen compound selected from nitrogen dioxide, nitrate or nitrite ions, nitrosylchloride, hydrazine or hydrazine compounds}</li> <li>{Separation; Purification}</li> <li>{Chlorine trioxide (ClO<sub>3</sub>); Chlorine hexoxide</li> </ul>
6/26 6/34 Halogens, co 7/00 7/01 7/012 7/015 7/017	<ul> <li>Preparation from the metal with the highest valency or from its oxides or salts of its oxyacids</li> <li>Purification; Stabilisation</li> <li>Purification; Stabilisation</li> <li>Purification; Stabilisation</li> <li>Padogens; Halogen acids (oxyacids C01B 11/00)</li> <li>Chlorine; Hydrogen chloride</li> <li>{Preparation of hydrogen chloride from the elements}</li> <li>{Chlorine hydrates; Obtaining chlorine therefrom}</li> <li>{Preparation of hydrogen chloride by reacting together chlorine, water and carbon or carbon monoxide (the carbon not acting only as catalyst)}</li> <li>Preparation from chlorides</li> <li>{Preparation of hydrogen chloride from chlorides}</li> <li>Preparation of chlorine from hydrogen chloride</li> <li>Preparation from ammonium chloride</li> </ul>	11/02 11/021 11/022 11/023 11/024 11/025 11/026 11/027	<ul> <li>Oxides of chlorine</li> <li>{Chlorine hemioxide (Cl<sub>2</sub>O)}</li> <li>{Chlorine dioxide (ClO<sub>2</sub>)}</li> <li>{Preparation from chlorites or chlorates}</li> <li>{from chlorites}</li> <li>{from chlorates without any other reaction reducing agent than chloride ions}</li> <li>{from chlorate ions in the presence of a peroxidic compound, e.g. hydrogen peroxide, ozone, peroxysulfates}</li> <li>{from chlorate ions in the presence of a nitrogen compound selected from nitrogen dioxide, nitrate or nitrite ions, nitrosylchloride, hydrazine or hydrazine compounds}</li> <li>{Separation; Purification}</li> <li>{Chlorine trioxide (ClO<sub>3</sub>); Chlorine hexoxide (Cl<sub>2</sub>O<sub>6</sub>); Chlorine heptoxide (Cl<sub>2</sub>O<sub>7</sub>)}</li> </ul>
6/26 6/34 Halogens, co 7/00 7/01 7/012 7/015 7/017	<ul> <li>Preparation from the metal with the highest valency or from its oxides or salts of its oxyacids</li> <li>Purification; Stabilisation</li> <li>Impounds thereof</li> <li>Halogens; Halogen acids (oxyacids C01B 11/00)</li> <li>Chlorine; Hydrogen chloride</li> <li>{Preparation of hydrogen chloride from the elements}</li> <li>{Chlorine hydrates; Obtaining chlorine therefrom}</li> <li>{Preparation of hydrogen chloride by reacting together chlorine, water and carbon or carbon monoxide (the carbon not acting only as catalyst)}</li> <li>Preparation from chlorides</li> <li>{Preparation of hydrogen chloride from chlorides}</li> <li>Preparation of chlorine from hydrogen chloride</li> <li>Preparation from ammonium chloride</li> <li>Preparation of hydrogen chloride from</li> </ul>	11/02 11/021 11/022 11/023 11/024 11/025 11/026 11/027	<ul> <li>Oxides of chlorine</li> <li>{Chlorine hemioxide (Cl<sub>2</sub>O)}</li> <li>{Chlorine dioxide (ClO<sub>2</sub>)}</li> <li>{Preparation from chlorites or chlorates}</li> <li>{from chlorites}</li> <li>{from chlorates without any other reaction reducing agent than chloride ions}</li> <li>{from chlorate ions in the presence of a peroxidic compound, e.g. hydrogen peroxide, ozone, peroxysulfates}</li> <li>{from chlorate ions in the presence of a nitrogen compound selected from nitrogen dioxide, nitrate or nitrite ions, nitrosylchloride, hydrazine or hydrazine compounds}</li> <li>{Separation; Purification}</li> <li>{Chlorine trioxide (ClO<sub>3</sub>); Chlorine hexoxide (Cl<sub>2</sub>O<sub>6</sub>); Chlorine heptoxide (Cl<sub>2</sub>O<sub>7</sub>)}</li> <li>Hypochlorous acid</li> </ul>
6/26 6/34 Halogens, co 7/00 7/01 7/012 7/015 7/017	<ul> <li>Preparation from the metal with the highest valency or from its oxides or salts of its oxyacids</li> <li>Purification; Stabilisation</li> <li>Purification; Stabilisation</li> <li>Purification; Stabilisation</li> <li>Padogens; Halogen acids (oxyacids C01B 11/00)</li> <li>Chlorine; Hydrogen chloride</li> <li>{Preparation of hydrogen chloride from the elements}</li> <li>{Chlorine hydrates; Obtaining chlorine therefrom}</li> <li>{Preparation of hydrogen chloride by reacting together chlorine, water and carbon or carbon monoxide (the carbon not acting only as catalyst)}</li> <li>Preparation from chlorides</li> <li>{Preparation of hydrogen chloride from chlorides}</li> <li>Preparation of chlorine from hydrogen chloride</li> <li>Preparation from ammonium chloride</li> </ul>	11/02 11/021 11/022 11/023 11/024 11/025 11/026 11/027	<ul> <li>Oxides of chlorine</li> <li>{Chlorine hemioxide (Cl<sub>2</sub>O)}</li> <li>{Chlorine dioxide (ClO<sub>2</sub>)}</li> <li>{Preparation from chlorites or chlorates}</li> <li>{from chlorites}</li> <li>{from chlorates without any other reaction reducing agent than chloride ions}</li> <li>{from chlorate ions in the presence of a peroxidic compound, e.g. hydrogen peroxide, ozone, peroxysulfates}</li> <li>{from chlorate ions in the presence of a nitrogen compound selected from nitrogen dioxide, nitrate or nitrite ions, nitrosylchloride, hydrazine or hydrazine compounds}</li> <li>{Separation; Purification}</li> <li>{Chlorine trioxide (ClO<sub>3</sub>); Chlorine hexoxide (Cl<sub>2</sub>O<sub>6</sub>); Chlorine heptoxide (Cl<sub>2</sub>O<sub>7</sub>)}</li> <li>Hypochlorous acid</li> <li>Hypochlorites</li> </ul>
6/26 6/34 Halogens, co 7/00 7/01 7/012 7/015 7/017	<ul> <li>Preparation from the metal with the highest valency or from its oxides or salts of its oxyacids</li> <li>Purification; Stabilisation</li> <li>Impounds thereof</li> <li>Halogens; Halogen acids (oxyacids C01B 11/00)</li> <li>Chlorine; Hydrogen chloride</li> <li>{Preparation of hydrogen chloride from the elements}</li> <li>{Chlorine hydrates; Obtaining chlorine therefrom}</li> <li>{Preparation of hydrogen chloride by reacting together chlorine, water and carbon or carbon monoxide (the carbon not acting only as catalyst)}</li> <li>Preparation from chlorides</li> <li>{Preparation of hydrogen chloride from chlorides}</li> <li>Preparation of chlorine from hydrogen chloride</li> <li>Preparation from ammonium chloride</li> <li>Preparation of hydrogen chloride from</li> </ul>	11/02 11/021 11/022 11/023 11/024 11/025 11/026 11/027 11/027 11/028 11/029 11/04 11/06 11/062	<ul> <li>Oxides of chlorine</li> <li>{Chlorine hemioxide (Cl<sub>2</sub>O)}</li> <li>{Chlorine dioxide (ClO<sub>2</sub>)}</li> <li>{Preparation from chlorites or chlorates}</li> <li>{from chlorites}</li> <li>{from chlorates without any other reaction reducing agent than chloride ions}</li> <li>{from chlorate ions in the presence of a peroxidic compound, e.g. hydrogen peroxide, ozone, peroxysulfates}</li> <li>{from chlorate ions in the presence of a nitrogen compound selected from nitrogen dioxide, nitrate or nitrite ions, nitrosylchloride, hydrazine or hydrazine compounds}</li> <li>{Separation; Purification}</li> <li>{Chlorine trioxide (ClO<sub>3</sub>); Chlorine hexoxide (Cl<sub>2</sub>O<sub>6</sub>); Chlorine heptoxide (Cl<sub>2</sub>O<sub>7</sub>)}</li> <li>Hypochlorous acid</li> <li>{Hypochlorites}</li> <li>{Hypochlorites of alkali metals}</li> </ul>
6/26 6/34 Halogens, co 7/00 7/01 7/012 7/015 7/017 7/03 7/035 7/04 7/05 7/055	<ul> <li>Preparation from the metal with the highest valency or from its oxides or salts of its oxyacids</li> <li>Purification; Stabilisation</li> <li>Impounds thereof</li> <li>Halogens; Halogen acids (oxyacids C01B 11/00)</li> <li>Chlorine; Hydrogen chloride</li> <li>{Preparation of hydrogen chloride from the elements}</li> <li>{Chlorine hydrates; Obtaining chlorine therefrom}</li> <li>{Preparation of hydrogen chloride by reacting together chlorine, water and carbon or carbon monoxide (the carbon not acting only as catalyst)}</li> <li>Preparation from chlorides</li> <li>{Preparation of hydrogen chloride from chlorides}</li> <li>Preparation of chlorine from hydrogen chloride</li> <li>Preparation from ammonium chloride</li> <li>Preparation of hydrogen chloride from ammonium chloride}</li> <li>Purification; {Separation (C01B 7/015 takes precedence)}</li> </ul>	11/02 11/021 11/022 11/023 11/024 11/025 11/026 11/027 11/027 11/028 11/029 11/04 11/06 11/062 11/064	<ul> <li>Oxides of chlorine</li> <li>{Chlorine hemioxide (Cl<sub>2</sub>O)}</li> <li>{Chlorine dioxide (ClO<sub>2</sub>)}</li> <li>{Preparation from chlorites or chlorates}</li> <li>{from chlorites}</li> <li>{from chlorates without any other reaction reducing agent than chloride ions}</li> <li>{from chlorate ions in the presence of a peroxidic compound, e.g. hydrogen peroxide, ozone, peroxysulfates}</li> <li>{from chlorate ions in the presence of a nitrogen compound selected from nitrogen dioxide, nitrate or nitrite ions, nitrosylchloride, hydrazine or hydrazine compounds}</li> <li>{Separation; Purification}</li> <li>{Chlorine trioxide (ClO<sub>3</sub>); Chlorine hexoxide (Cl<sub>2</sub>O<sub>6</sub>); Chlorine heptoxide (Cl<sub>2</sub>O<sub>7</sub>)}</li> <li>Hypochlorites</li> <li>{Hypochlorites of alkali metals}</li> <li>{Hypochlorites of alkaline-earth metals}</li> </ul>
6/26 6/34 Halogens, co 7/00 7/01 7/012 7/015 7/017 7/03 7/035 7/04 7/05 7/055	<ul> <li>Preparation from the metal with the highest valency or from its oxides or salts of its oxyacids</li> <li>Purification; Stabilisation</li> <li>Impounds thereof</li> <li>Halogens; Halogen acids (oxyacids C01B 11/00)</li> <li>Chlorine; Hydrogen chloride</li> <li>{Preparation of hydrogen chloride from the elements}</li> <li>{Chlorine hydrates; Obtaining chlorine therefrom}</li> <li>{Preparation of hydrogen chloride by reacting together chlorine, water and carbon or carbon monoxide (the carbon not acting only as catalyst)}</li> <li>Preparation from chlorides</li> <li>{Preparation of hydrogen chloride from chlorides}</li> <li>Preparation of chlorine from hydrogen chloride</li> <li>Preparation from ammonium chloride</li> <li>Preparation of hydrogen chloride from ammonium chloride}</li> <li>Purification; {Separation (C01B 7/015 takes)</li> </ul>	11/02 11/021 11/022 11/023 11/024 11/025 11/026 11/027 11/027 11/028 11/029 11/04 11/06 11/062 11/064 11/066	<ul> <li>Oxides of chlorine</li> <li>{Chlorine hemioxide (Cl<sub>2</sub>O)}</li> <li>{Chlorine dioxide (ClO<sub>2</sub>)}</li> <li>{Preparation from chlorites or chlorates}</li> <li>{from chlorites}</li> <li>{from chlorates without any other reaction reducing agent than chloride ions}</li> <li>{from chlorate ions in the presence of a peroxidic compound, e.g. hydrogen peroxide, ozone, peroxysulfates}</li> <li>{from chlorate ions in the presence of a nitrogen compound selected from nitrogen dioxide, nitrate or nitrite ions, nitrosylchloride, hydrazine or hydrazine compounds}</li> <li>{Separation; Purification}</li> <li>{Chlorine trioxide (ClO<sub>3</sub>); Chlorine hexoxide (Cl<sub>2</sub>O<sub>6</sub>); Chlorine heptoxide (Cl<sub>2</sub>O<sub>7</sub>)}</li> <li>Hypochlorites</li> <li>{Hypochlorites of alkali metals}</li> <li>{Magnesium hypochlorite}</li> </ul>
6/26 6/34 Halogens, co 7/00 7/01 7/012 7/015 7/017 7/03 7/035 7/04 7/05 7/055 7/07	<ul> <li>Preparation from the metal with the highest valency or from its oxides or salts of its oxyacids</li> <li>Purification; Stabilisation</li> <li>Impounds thereof</li> <li>Halogens; Halogen acids (oxyacids C01B 11/00)</li> <li>Chlorine; Hydrogen chloride</li> <li>{Preparation of hydrogen chloride from the elements}</li> <li>{Chlorine hydrates; Obtaining chlorine therefrom}</li> <li>{Preparation of hydrogen chloride by reacting together chlorine, water and carbon or carbon monoxide (the carbon not acting only as catalyst)}</li> <li>Preparation from chlorides</li> <li>{Preparation of hydrogen chloride from chlorides}</li> <li>Preparation of chlorine from hydrogen chloride</li> <li>Preparation from ammonium chloride</li> <li>Preparation of hydrogen chloride from ammonium chloride}</li> <li>Purification; {Separation (C01B 7/015 takes precedence)}</li> </ul>	11/02 11/021 11/022 11/023 11/024 11/025 11/026 11/027 11/027 11/028 11/029 11/04 11/06 11/062 11/064 11/066	<ul> <li>Oxides of chlorine</li> <li>{Chlorine hemioxide (Cl<sub>2</sub>O)}</li> <li>{Chlorine dioxide (ClO<sub>2</sub>)}</li> <li>{Preparation from chlorites or chlorates}</li> <li>{from chlorites}</li> <li>{from chlorates without any other reaction reducing agent than chloride ions}</li> <li>{from chlorate ions in the presence of a peroxidic compound, e.g. hydrogen peroxide, ozone, peroxysulfates}</li> <li>{from chlorate ions in the presence of a nitrogen compound selected from nitrogen dioxide, nitrate or nitrite ions, nitrosylchloride, hydrazine or hydrazine compounds}</li> <li>{Separation; Purification}</li> <li>{Chlorine trioxide (ClO<sub>3</sub>); Chlorine hexoxide (Cl<sub>2</sub>O<sub>6</sub>); Chlorine heptoxide (Cl<sub>2</sub>O<sub>7</sub>)}</li> <li>Hypochlorites</li> <li>{Hypochlorites of alkali metals}</li> <li>{Magnesium hypochlorite}</li> <li>{Stabilisation by additives other than oxides,</li> </ul>
6/26 6/34 Halogens, co 7/00 7/01 7/012 7/015 7/017 7/03 7/035 7/04 7/05 7/055 7/07	<ul> <li>Preparation from the metal with the highest valency or from its oxides or salts of its oxyacids</li> <li>Purification; Stabilisation</li> <li>Impounds thereof</li> <li>Halogens; Halogen acids (oxyacids C01B 11/00)</li> <li>Chlorine; Hydrogen chloride</li> <li>{Preparation of hydrogen chloride from the elements}</li> <li>{Chlorine hydrates; Obtaining chlorine therefrom}</li> <li>{Preparation of hydrogen chloride by reacting together chlorine, water and carbon or carbon monoxide (the carbon not acting only as catalyst)}</li> <li>Preparation from chlorides</li> <li>{Preparation of hydrogen chloride from chlorides}</li> <li>Preparation of chlorine from hydrogen chloride</li> <li>Preparation from ammonium chloride</li> <li>Preparation (C01B 7/015 takes precedence)}</li> <li>{of hydrogen chloride}</li> </ul>	11/02 11/021 11/022 11/023 11/024 11/025 11/026 11/027 11/027 11/028 11/029 11/04 11/06 11/062 11/064 11/066	<ul> <li>Oxides of chlorine</li> <li>{Chlorine hemioxide (Cl<sub>2</sub>O)}</li> <li>{Chlorine dioxide (ClO<sub>2</sub>)}</li> <li>{Preparation from chlorites or chlorates}</li> <li>{from chlorites}</li> <li>{from chlorates without any other reaction reducing agent than chloride ions}</li> <li>{from chlorate ions in the presence of a peroxidic compound, e.g. hydrogen peroxide, ozone, peroxysulfates}</li> <li>{from chlorate ions in the presence of a nitrogen compound selected from nitrogen dioxide, nitrate or nitrite ions, nitrosylchloride, hydrazine or hydrazine compounds}</li> <li>{Separation; Purification}</li> <li>{Chlorine trioxide (ClO<sub>3</sub>); Chlorine hexoxide (Cl<sub>2</sub>O<sub>6</sub>); Chlorine heptoxide (Cl<sub>2</sub>O<sub>7</sub>)}</li> <li>Hypochlorous acid</li> <li>Hypochlorites</li> <li>{Hypochlorites of alkali metals}</li> <li>{Magnesium hypochlorite}</li> <li>{Stabilisation by additives other than oxides, hydroxides, carbonates of alkali or alkaline-earth metals; Coating of particles; Shaping; Granulation}</li> </ul>
6/26 6/34 Halogens, co 7/00 7/01 7/012 7/015 7/017 7/03 7/035 7/04 7/05 7/055 7/07 7/0706 7/0712	<ul> <li>Preparation from the metal with the highest valency or from its oxides or salts of its oxyacids</li> <li>Purification; Stabilisation</li> <li>Purification; Stabilisation</li> <li>Purification; Stabilisation</li> <li>Preparation of the coloride (oxyacids C01B 11/00)</li> <li>Chlorine; Hydrogen chloride</li> <li>{Preparation of hydrogen chloride from the elements}</li> <li>{Chlorine hydrates; Obtaining chlorine therefrom}</li> <li>{Preparation of hydrogen chloride by reacting together chlorine, water and carbon or carbon monoxide (the carbon not acting only as catalyst)}</li> <li>Preparation from chlorides</li> <li>{Preparation of hydrogen chloride from chlorides}</li> <li>Preparation of chlorine from hydrogen chloride</li> <li>Preparation from ammonium chloride</li> <li>Preparation; {Separation (C01B 7/015 takes precedence)}</li> <li>{of hydrogen chloride}</li> <li>{by distillation}</li> </ul>	11/02 11/021 11/022 11/023 11/024 11/025 11/026 11/027 11/027 11/028 11/029 11/04 11/06 11/062 11/064 11/066	<ul> <li>Oxides of chlorine</li> <li>{Chlorine hemioxide (Cl<sub>2</sub>O)}</li> <li>{Chlorine dioxide (ClO<sub>2</sub>)}</li> <li>{Preparation from chlorites or chlorates}</li> <li>{from chlorites}</li> <li>{from chlorates without any other reaction reducing agent than chloride ions}</li> <li>{from chlorate ions in the presence of a peroxidic compound, e.g. hydrogen peroxide, ozone, peroxysulfates}</li> <li>{from chlorate ions in the presence of a nitrogen compound selected from nitrogen dioxide, nitrate or nitrite ions, nitrosylchloride, hydrazine or hydrazine compounds}</li> <li>{Separation; Purification}</li> <li>{Chlorine trioxide (ClO<sub>3</sub>); Chlorine hexoxide (Cl<sub>2</sub>O<sub>6</sub>); Chlorine heptoxide (Cl<sub>2</sub>O<sub>7</sub>)}</li> <li>Hypochlorous acid</li> <li>Hypochlorites</li> <li>{Hypochlorites of alkali metals}</li> <li>{Magnesium hypochlorite}</li> <li>{Stabilisation by additives other than oxides, hydroxides, carbonates of alkali or alkaline-earth metals; Coating of particles; Shaping;</li> </ul>
6/26 6/34 Halogens, co 7/00 7/01 7/012 7/015 7/017 7/03 7/035 7/04 7/05 7/055 7/07 7/0706 7/0712 7/0718	<ul> <li>Preparation from the metal with the highest valency or from its oxides or salts of its oxyacids</li> <li>Purification; Stabilisation</li> <li>Impounds thereof</li> <li>Halogens; Halogen acids (oxyacids C01B 11/00)</li> <li>Chlorine; Hydrogen chloride</li> <li>{Preparation of hydrogen chloride from the elements}</li> <li>{Chlorine hydrates; Obtaining chlorine therefrom}</li> <li>{Preparation of hydrogen chloride by reacting together chlorine, water and carbon or carbon monoxide (the carbon not acting only as catalyst)}</li> <li>Preparation from chlorides</li> <li>{Preparation of hydrogen chloride from chlorides}</li> <li>Preparation of chlorine from hydrogen chloride</li> <li>Preparation from ammonium chloride</li> <li>Preparation (C01B 7/015 takes precedence)}</li> <li>{of hydrogen chloride}</li> <li>{by distillation}</li> <li>{by dasorption}</li> </ul>	11/02 11/021 11/022 11/023 11/024 11/025 11/026 11/027 11/028 11/029 11/04 11/06 11/062 11/064 11/068 11/068	<ul> <li>Oxides of chlorine</li> <li>{Chlorine hemioxide (Cl<sub>2</sub>O)}</li> <li>{Chlorine dioxide (ClO<sub>2</sub>)}</li> <li>{Preparation from chlorites or chlorates}</li> <li>{from chlorites}</li> <li>{from chlorates without any other reaction reducing agent than chloride ions}</li> <li>{from chlorate ions in the presence of a peroxidic compound, e.g. hydrogen peroxide, ozone, peroxysulfates}</li> <li>{from chlorate ions in the presence of a nitrogen compound selected from nitrogen dioxide, nitrate or nitrite ions, nitrosylchloride, hydrazine or hydrazine compounds}</li> <li>{Separation; Purification}</li> <li>{Chlorine trioxide (ClO<sub>3</sub>); Chlorine hexoxide (Cl<sub>2</sub>O<sub>6</sub>); Chlorine heptoxide (Cl<sub>2</sub>O<sub>7</sub>)}</li> <li>Hypochlorous acid</li> <li>Hypochlorites</li> <li>{Hypochlorites of alkali metals}</li> <li>{Magnesium hypochlorite}</li> <li>{Stabilisation by additives other than oxides, hydroxides, carbonates of alkali or alkalineearth metals; Coating of particles; Shaping; Granulation}</li> <li>Chlorous acid</li> <li>Chlorites</li> </ul>
6/26 6/34 Halogens, co 7/00 7/01 7/012 7/015 7/017 7/03 7/035 7/04 7/05 7/055 7/07 7/0706 7/0712 7/0718 7/0725	<ul> <li>Preparation from the metal with the highest valency or from its oxides or salts of its oxyacids</li> <li>Purification; Stabilisation</li> <li>Impounds thereof</li> <li>Halogens; Halogen acids (oxyacids C01B 11/00)</li> <li>Chlorine; Hydrogen chloride</li> <li>{Preparation of hydrogen chloride from the elements}</li> <li>{Chlorine hydrates; Obtaining chlorine therefrom}</li> <li>{Preparation of hydrogen chloride by reacting together chlorine, water and carbon or carbon monoxide (the carbon not acting only as catalyst)}</li> <li>Preparation from chlorides</li> <li>{Preparation of hydrogen chloride from chlorides}</li> <li>Preparation of chlorine from hydrogen chloride</li> <li>Preparation from ammonium chloride</li> <li>Preparation (C01B 7/015 takes precedence)}</li> <li>{of hydrogen chloride}</li> <li>{by distillation}</li> <li>{by dasorption}</li> <li>{by active carbon}</li> </ul>	11/02 11/021 11/022 11/023 11/024 11/025 11/026 11/027 11/028 11/029 11/04 11/06 11/062 11/064 11/066 11/068	<ul> <li>Oxides of chlorine</li> <li>{Chlorine hemioxide (Cl<sub>2</sub>O)}</li> <li>{Chlorine dioxide (ClO<sub>2</sub>)}</li> <li>{Preparation from chlorites or chlorates}</li> <li>{from chlorites}</li> <li>{from chlorates without any other reaction reducing agent than chloride ions}</li> <li>{from chlorate ions in the presence of a peroxidic compound, e.g. hydrogen peroxide, ozone, peroxysulfates}</li> <li>{from chlorate ions in the presence of a nitrogen compound selected from nitrogen dioxide, nitrate or nitrite ions, nitrosylchloride, hydrazine or hydrazine compounds}</li> <li>{Separation; Purification}</li> <li>{Chlorine trioxide (ClO<sub>3</sub>); Chlorine hexoxide (Cl<sub>2</sub>O<sub>6</sub>); Chlorine heptoxide (Cl<sub>2</sub>O<sub>7</sub>)}</li> <li>Hypochlorous acid</li> <li>Hypochlorites</li> <li>{Hypochlorites of alkali metals}</li> <li>{Magnesium hypochlorite}</li> <li>{Stabilisation by additives other than oxides, hydroxides, carbonates of alkali or alkaline-earth metals; Coating of particles; Shaping; Granulation}</li> <li>Chlorous acid</li> </ul>

11/14	Chlorates	13/10	Preparation of ozone
11/145	• • • {Separation; Crystallisation; Purification,	13/11	by electric discharge
	After-treatment; Stabilisation by additives}		NOTE
11/16	Perchloric acid		11012
11/18	Perchlorates		In groups <u>C01B 13/11</u> and <u>C01B 13/115</u> ,
11/185	{Ammonium perchlorate}		additional features relating to the
11/20	Oxygen compounds of bromine		preparation of ozone by electrical discharge
11/22	Oxygen compounds of iodine		are indexed with codes chosen from C01B 2201/00 - C01B 2201/90.
11/24	. Oxygen compounds of fluorine		
Oxygen: Oxi	des or hydroxides in general; Per-compounds	13/115	• • • {characterised by the electrical circuits
		13/14	producing the electrical discharge}  Methods for preparing oxides or hydroxides in
13/00	Oxygen; Ozone; Oxides or hydroxides in general	13/14	general (particular individual oxides or hydroxides,
13/02	• Preparation of oxygen (by liquefying <u>F25J</u> )		see the relevant groups of subclasses C01B - C01G
13/0203	• • {from inorganic compounds}		or C25B, according to the element combined with
13/0207	{Water}		the oxygen or hydroxy group)
13/0211 13/0214	<ul><li> {Peroxy compounds}</li><li> {Hydrogen peroxide}</li></ul>	13/145	• • {After-treatment of oxides or hydroxides, e.g.
13/0214	{Chlorate}		pulverising, drying, decreasing the acidity}
13/0218	{from organic compounds}	13/16	Purification
13/0225	{Peroxy compounds}	13/18	• by thermal decomposition of compounds, e.g. of
13/0229	{Purification or separation processes}	12/105	salts or hydroxides
13/022)		13/185	• • {Preparing mixtures of oxides}
	<u>NOTE</u>	13/20	<ul> <li>by oxidation of elements in the gaseous state;</li> <li>by oxidation or hydrolysis of compounds in the</li> </ul>
	In groups <u>C01B 13/0229</u> - <u>C01B 13/0288</u> ,		gaseous state
	additional features relating to the	13/22	of halides or oxyhalides
	purification or separation processes	13/24	in the presence of hot combustion gases
	are indexed with codes chosen from	13/26	in the presence of a fluidised bed
	<u>C01B 2210/0026</u> - <u>C01B 2210/0098</u> .	13/28	using a plasma or an electric discharge
13/0233	• • • {Chemical processing only}	13/30	Removal and cooling of the oxide-containing
13/0237	• • • {by oxidation}		suspension
13/024	• • • {by reduction}	13/32	• • by oxidation or hydrolysis of elements or
13/0244	• • • {by complexation}		compounds in the liquid or solid state {or in non-
13/0248	• • • {Physical processing only}	10/000	aqueous solution, e.g. sol-gel process}
13/0251	• • • {by making use of membranes}	13/322	• • • {of elements or compounds in the solid state}
13/0255	• • • • {characterised by the type of membrane}	13/324	• • • {by solid combustion synthesis}
13/0259	• • • {by adsorption on solids}	13/326	• • • {of elements or compounds in the liquid state}
13/0262	{characterised by the adsorbent}	13/328	• • {by processes making use of emulsions, e.g. the kerosine process}
13/0266	{Carbon based materials}	13/34	<ul> <li>by oxidation or hydrolysis of sprayed or atomised</li> </ul>
13/027	{Zeolites}	13/31	solutions
13/0274	{Other molecular sieve materials}	13/36	• • by precipitation reactions in {aqueous} solutions
13/0277 13/0281	{Temperature swing adsorption}		$\{(\underline{\text{C01B } 13/328} \text{ takes precedence})\}$
13/0281	<ul><li> {in getters}</li><li> {by absorption in liquids}</li></ul>	13/363	• • • {Mixtures of oxides or hydroxides by
13/0288	{Combined chemical and physical processing}		precipitation}
13/0200		13/366	• • • {by hydrothermal processing}
	NOTE	15/00	Peroxides; Peroxyhydrates; Peroxyacids or salts
	In this group, processing steps are		thereof; Superoxides; Ozonides
	indexed with codes chosen from	15/005	• {Stabilisation of the solid compounds subsequent to
	<u>C01B 2210/0001</u> - <u>C01B 2210/0025</u>		the preparation or to the crystallisation, by additives
13/0292	• • {Preparation from air using a molten phase		or by coating}
	containing alkali metal nitrite, optionally together	15/01	. Hydrogen peroxide
	with other oxygen acceptors}	15/013	• Separation; Purification; Concentration
13/0296	• • {Generators releasing in a self-sustaining	15/0135	• • (Purification by solid ion-exchangers or solid
	way pure oxygen from a solid charge, without	15/017	chelating agents} Anhydrous hydrogen peroxide; Anhydrous
	interaction of it with a fluid nor external heating,	13/01/	solutions or gaseous mixtures containing
	e.g. chlorate candles or canisters containing them		hydrogen peroxide
13/08	<ul> <li>(charges per se C01B 13/02)}</li> <li>from air with the aid of metal oxides, e.g. barium</li> </ul>	15/022	Preparation from organic compounds
13/00	oxide, manganese oxide {(C01B 13/0292 takes	15/023	by the alkyl-anthraquinone process
	precedence)}	15/024	• • • from hydrocarbons
13/083	• • {with barium oxide}	15/026	from alcohols
13/086	{with manganese oxide}	15/027	• • Preparation from water

15/0275	{Preparation by reaction of water, carbon	17/0408 {Pretreatment of the hydrogen sulfide
15/020	monoxide and oxygen}	containing gases}
15/029	• Preparation from hydrogen and oxygen	17/0413 {characterised by the combustion step}
15/0295	• • {by electrical discharge}	17/0417 {Combustion reactors}
15/03	Preparation from inorganic peroxy compounds,	17/0421 {Multistage combustion}
1.7/0.00	e.g. from peroxysulfate	17/0426 {characterised by the catalytic conversion}
15/032	from metal peroxides	17/043 {Catalytic converters}
15/037	Stabilisation by additives	17/0434 {Catalyst compositions}
15/04	• Metal peroxides or peroxyhydrates thereof; {Metal}	17/0439 {at least one catalyst bed operating below
	superoxides; {Metal} ozonides; {Peroxyhydrates	the dew-point of sulfur}
	thereof}	17/0443 {in a moving bed}
15/043	of alkali metals, alkaline earth metals or	17/0447 {Separation of the obtained sulfur}
	magnesium {or beryllium or aluminium}	17/0452 {Process control; Start-up or cooling-down
15/0435	• • · {of alkali metals}	procedures of the Claus process}
15/047	of heavy metals	17/0456 {the hydrogen sulfide-containing gas being a
15/0475	• • { of actinides }	Claus process tail gas}
15/055	• Peroxyhydrates ( <u>C01B 15/04</u> takes precedence);	17/046 { without intermediate formation of sulfur
	Peroxyacids or salts thereof	dioxide}
15/06	containing sulfur	17/0465 {Catalyst compositions}
15/08	Peroxysulfates	17/0469 {at least one catalyst bed operating below
15/085	{Stabilisation of the solid compounds,	the dew-point of sulfur}
	subsequent to the preparation or to the	17/0473 {by reaction of sulfur dioxide or sulfur trioxide
	crystallisation, by additives or by coating}	containing gases with reducing agents other
15/10	containing carbon	than hydrogen sulfide}
15/103	{containing only alkali metals as metals}	17/0478 { with hydrocarbons or mixtures containing
15/106	• • • {Stabilisation of the solid compounds,	them}
	subsequent to the preparation or to the	17/0482 { with carbon or solid carbonaceous
	crystallisation, by additives or by coating}	materials}
15/12	• containing boron	17/0486 { with carbon monoxide or carbon monoxide
15/123	{Stabilisation of the solid compounds,	containing mixtures}
	subsequent to the preparation or to the	17/0491 { with hydrogen or hydrogen-containing
	crystallisation, by additives or by coating}	mixtures, e.g. synthesis gas}
15/126	{Dehydration of solid hydrated peroxyborates	17/0495 {by dissociation of hydrogen sulfide into the
	to less hydrated or anhydrous products}	elements}
15/14	• containing silicon	17/05 by wet processes
15/16	containing phosphorus	17/06 from non-gaseous sulfides or materials containing
		such sulfides, e.g. ores
17/00	Sulfur; Compounds thereof	17/10 . Finely divided sulfur, e.g. sublimed sulfur,
17/02	<ul> <li>Preparation of sulfur; Purification</li> </ul>	flowers of sulfur
17/0205	• • {Separation of sulfur from liquids, e.g. by	17/12 Insoluble sulfur (mu-sulfur)
	coalescence}	17/125 {Sulfur isotopes other than 32S}
17/021	• • {Separation of sulfur from gases}	17/16 • Hydrogen sulfides
17/0216	• • {Solidification or cooling of liquid sulfur}	17/161 {Preparation from elemental sulfur}
17/0221	• • {Melting}	17/162 {from elemental sulfur and hydrogen}
17/0226	• • {Vaporising or superheating}	17/164 {Preparation by reduction of oxidic sulfur
17/0232	• • {Purification, e.g. degassing}	compounds}
17/0237	• • {Converting into particles, e.g. by granulation,	17/165 • • {Preparation from sulfides, oxysulfides or
	milling}	polysulfides}
17/0243	• • {Other after-treatment of sulfur}	17/167 • • {Separation}
17/0248	• • • {of particulate sulfur}	17/168 . {Purification}
17/0253	• • {from non-gaseous sulfur compounds other than	17/18 Hydrogen polysulfides
	sulfides or materials containing such sulfides}	17/20 • Methods for preparing sulfides or polysulfides,
17/0259	• • • {by reduction of sulfates}	in general (ammonium sulfides or polysulfides
17/0264	• • • {of calcium sulfates}	<u>C01C</u> ; sulfides or polysulfides of metals, other than
17/027	Recovery of sulfur from material containing	alkali metals, magnesium, calcium, strontium and
	elemental sulfur, e.g. luxmasses {or sulfur	barium, <u>see</u> the relevant groups of subclasses <u>C01F</u>
	containing ores); Purification (of the recovered	or <u>C01G</u> , according to the metal)
	sulfur}	17/22 • Alkali metal sulfides or polysulfides
17/033	using a liquid extractant	17/24 • Preparation by reduction
17/04	from gaseous sulfur compounds including	17/26 with carbon
	gaseous sulfides	17/28 with reducing gases
17/0404	• • • {by processes comprising a dry catalytic	17/30 With reducing gases  17/30 Preparation from sodium or potassium amalgam
	conversion of hydrogen sulfide-containing	with sulfur or sulfides
	gases, e.g. the Claus process}	with sulful of sulfides

17/32	Hydrosulfides of sodium or potassium	17/76	hy contact processes
	Polysulfides of sodium or potassium		by contact processes
17/34		17/762	{High pressure processes}
17/36	. Purification	17/765	Multi-stage SO <sub>3</sub> -conversion
17/38	. Dehydration	17/7655	• • • • { with intermediate absorption }
17/40	. Making shaped products, e.g. granules	17/77	Fluidised-bed processes
17/42	<ul> <li>Sulfides or polysulfides of magnesium, calcium, strontium, or barium</li> </ul>	17/775	Liquid phase contacting processes or wet catalysis processes
17/43	<ul> <li>from oxides or hydroxides with sulfur or</li> </ul>	17/78	characterised by the catalyst used
	hydrogen sulfide	17/79	containing vanadium
17/44	• • by reduction of sulfates	17/80	Apparatus
17/45	<ul> <li>Compounds containing sulfur and halogen, with or</li> </ul>	17/803	• • • • {Converters}
	without oxygen	17/806	{Absorbers; Heat exchangers}
17/4507	• • {containing sulfur and halogen only}	17/82	of sulfuric acid using a nitrogen oxide process
17/4515	• • {containing sulfur and fluorine only}	17/84	Chamber process
17/4523	• • • {Sulfur tetrafluoride}	17/86	Tower process
17/453	• • • {Sulfur hexafluoride}	17/88	Concentration of sulfuric acid
17/4538	• • {containing sulfur and chlorine only}	17/90	Separation; Purification
17/4546	• • • {Sulfur dichloride}	17/901	• • • {Recovery from spent acids containing metallic
17/4553	• • • {Sulfur hexachloride}		ions, e.g. hydrolysis acids, pickling acids
17/4561	{Compounds containing sulfur, halogen and		(obtaining sulfur dioxide as an intermediate
	oxygen only}		in sulfur trioxide recovery from sulfates, e.g.
17/4569	• • • {Thionyl fluoride (SOF <sub>2</sub> )}		iron sulfates <u>C01B 17/501</u> , from spent acids
17/4576	• • • {Sulfuryl fluoride $(SO_2F_2)$ }		<u>C01B 17/58</u> )}
17/4584	• • • {Thionyl chloride (SOCl <sub>2</sub> )}	17/902	• • • {by dialysis}
17/4592	• • • {Sulfuryl chloride (SO <sub>2</sub> Cl <sub>2</sub> )}	17/903	• • • {by liquid-liquid extraction}
17/46	<ul> <li>Compounds containing sulfur, halogen, hydrogen,</li> </ul>	17/904	• • • {by ion-exchange}
	and oxygen	17/905	{Removal of organic impurities}
17/463	• • {Fluorosulfonic acid (FSO <sub>3</sub> H)}	17/906	{Removal of mercury}
17/466	• • {Chlorosulfonic acid (ClSO <sub>3</sub> H)}	17/907	{Removal of arsenic}
17/48	Sulfur dioxide; Sulfurous acid	17/908	• • • {Removal of antimony or bismuth}
17/50	Preparation of sulfur dioxide	17/92	Recovery from acid tar or the like, {e.g.
17/501	• • {by reduction of sulfur compounds}		alkylation acids (obtaining sulfur dioxide as
17/502	{of sulfur trioxide}		an intermediate in sulfur trioxode recovery
17/503	{of sulfuric acid}	17/025	therefrom <u>C01B 17/58</u> )}
17/504	• • • {of ammonium sulfates (of ammonium	17/925	• • • • {by processes involving a liquid-liquid extraction}
	sulfates containing sulfuric acid solutions	17/94	Recovery from nitration acids
17/505	<u>C01B 17/585</u> )}	17/96	• Methods for the preparation of sulfates in general
17/505	{of alkali metal sulfates}		(particular individual sulfates, see the relevant
17/506	• • • {of calcium sulfates}		groups of subclasses <u>C01B</u> - <u>C01G</u> , according to the
17/507	{of iron sulfates}		cation)
17/508	• • • {by oxidation of sulfur compounds}	17/965	• • {Pyrosulfates}
17/52	• • • by roasting sulfides ( <u>C22B 1/00</u> takes	17/98	Other compounds containing sulfur and oxygen
17/5/	precedence)		(persulfuric acids <u>C01B 15/06</u> ; persulfates
17/54 17/56	<ul><li> by burning elemental sulfur</li><li> Separation; Purification</li></ul>		<u>C01B 15/08</u> )
	-	19/00	Selenium; Tellurium; Compounds thereof
17/58	Recovery of sulfur dioxide from acid tar or the like {or from any waste sulfuric acid}		(phosphorus compounds <u>C01B 25/14</u> )
17/585	{from ammonium sulfate containing sulfuric acid solutions}	19/001	• {Preparation involving a liquid-liquid extraction, an adsorption or an ion-exchange}
17/60	Isolation of sulfur dioxide from gases	19/002	• {Compounds containing, besides selenium or
17/62	Methods of preparing sulfites in general (particular)		tellurium, more than one other element, with -O-
17702	individual sulfites, see the relevant groups of		and -OH not being considered as anions}
	subclasses C01B - C01G, according to the cation)	19/004	• {Oxides; Hydroxides}
17/625	• {metabisulfites or pyrosulfites}	19/005	• {Halides}
17/64	• Thiosulfates; Dithionites; Polythionates	19/007	• {Tellurides or selenides of metals ( <u>C01B 19/002</u>
17/66	• Dithionites {or hydrosulfites (S <sub>2</sub> O <sub>4</sub> <sup>2-</sup> )}		takes precedence)}
17/665	• • Stabilisation by additives subsequent to	19/008	• {Salts of oxyacids of selenium or tellurium}
17/003	preparation; Dust prevention by additives}	19/02	Elemental selenium or tellurium
17/69	Sulfur trioxide; Sulfuric acid	19/04	Binary compounds {including binary selenium-
17/70	Stabilisation of gamma-form sulfur trioxide		tellurium compounds ( <u>C01B 19/004</u> , <u>C01B 19/005</u> ,
17/74	Preparation		C01B 19/007 take precedence)
17/745	{from sulfates}	21/00	Nitrogen; Compounds thereof
	(		

21/02	Dramaration of nitro can (by decomposition of	21/0625 (with compan)
21/02	<ul> <li>Preparation of nitrogen (by decomposition of ammonia {C01B 3/047})</li> </ul>	21/0625 {with copper}
21/04	<ul> <li>Purification or separation of nitrogen (by liquefying</li> </ul>	21/0627 {with one or more rare earth metals}
21/04	F25J)	21/063 {with one or more actinides, e.g. UN, PuN}
21/0405	• • {Purification or separation processes}	21/0632 • { with gallium, indium or thallium}
21/0403	• • {1 utilication of separation processes}	21/0635 • • {with germanium, tin or lead}
	<u>NOTE</u>	21/0637 {with metals not specified in groups
	In this group, additional features relating	C01B 21/0607 - C01B 21/0635, other than aluminium, titanium, zirconium or hafnium}
	to the purification or separation processes	
	are indexed with codes chosen from	21/064 with boron
	C01B 2210/0026 - C01B 2210/0098	21/0641 {Preparation by direct nitridation of elemental boron}
01/0411		•
21/0411	• • • {Chemical processing only}	,
21/0416	• • • {by oxidation}	
21/0422	• • • {by reduction}	21/0646 {Preparation by pyrolysis of boron and nitrogen containing compounds}
21/0427	• • • {by complexation}	
21/0433	• • • {Physical processing only}	21/0648 {After-treatment, e.g. grinding, purification (transformation of hexagonal into cubic or
21/0438	• • • {by making use of membranes}	wurtzitic boron nitride C04B 35/5831)}
21/0444	• • • • {characterised by the membrane}	21/068 • with silicon
21/045	• • • {by adsorption in solids}	21/0682 {Preparation by direct nitridation of silicon}
21/0455	• • • • {characterised by the adsorbent}	21/0685 {Preparation by direct intriduction of sincon}
21/0461	{Carbon based materials}	* * *
21/0466	• • • • • {Zeolites}	( , , , , , , , , , , , , , , , , , , ,
21/0472	• • • • • {Other molecular sieve materials}	
21/0477	• • • • {Temperature swing adsorption}	21/0722 • • • {Preparation by direct nitridation of aluminium}
21/0483	{in getters}	
21/0488	• • • {by absorption in liquids}	21/0724 {using a plasma}
21/0494	• • {Combined chemical and physical processing}	21/0726 {Preparation by carboreductive nitridation}
	NOTE	21/0728 {After-treatment, e.g. grinding, purification}
	NOTE	21/076 • with titanium or zirconium {or hafnium}
	In this group, processing steps are	21/0761 {Preparation by direct nitridation of titanium, zirconium or hafnium}
	indexed with codes chosen from	,
	<u>C01B 2210/0001</u> - <u>C01B 2210/0025</u>	21/0763 {Preparation from titanium, zirconium or hafnium halides}
21/06	. Binary compounds of nitrogen with metals, with	21/0765 {Preparation by carboreductive nitridation}
	silicon, or with boron, {or with carbon, i.e. nitrides;	21/0766 {Preparation by pyrolysis of nitrogen
	Compounds of nitrogen with more than one metal,	containing titanium, zirconium or hafnium
	silicon or boron} (azides C01B 21/08)	compounds}
	<u>NOTES</u>	21/0768 • • • {After-treatment, e.g. grinding, purification}
	110 120	21/08 • Hydrazoic acid; Azides; Halogen azides
	1. Binary compounds, i.e. compounds of nitrogen	21/082 • Compounds containing nitrogen and non-metals
	with only one other element chosen from metals,	{and optionally metals} (C01B 21/06, C01B 21/08
	silicon, boron or carbon, are classified in groups	take precedence)
	<u>C01B 21/06</u> or <u>C01B 21/0605</u> - <u>C01B 21/076</u> .	21/0821 • • {Oxynitrides of metals, boron or silicon}
	Compounds of nitrogen with more than one	21/0823 {Silicon oxynitrides}
	element chosen from metals, silicon or boron are	21/0825 {Aluminium oxynitrides}
	classified in <u>C01B 21/0602</u>	21/0826 {Silicon aluminium oxynitrides, i.e. sialons}
	2. Documents relating to several specific binary	21/0828 • • {Carbonitrides or oxycarbonitrides of metals,
	compounds are classified in C01B 21/06 only	boron or silicon}
	and receive the indexing codes chosen from	21/083 containing one or more halogen atoms
	$\frac{\text{C01B } 21/0602}{\text{C01B } 21/076}$ to identify the	21/0832 {Binary compounds of nitrogen with halogens}
	specific compounds	21/0835 {Nitrogen trifluoride}
21/0602	• • { with two or more other elements chosen from	21/0837 {Purification}
	metals, silicon or boron}	21/084 containing also one or more oxygen atoms, e.g.
21/0605	• • {Binary compounds of nitrogen with carbon}	nitrosyl halides
21/0607	• • {with alkali metals}	21/0842 {Halides of nitrogen oxides}
21/061	• • {with lithium}	21/0844 {Nitrosyl fluoride}
21/0612	• • {with alkaline-earth metals, beryllium or	21/0846 {Nitrosyl chloride}
	magnesium}	21/0848 {Nitrosyl perchlorate}
21/0615		
21/0013	• • {with transition metals other than titanium,	21/086 • • containing one or more sulfur atoms
	zirconium or hafnium}	21/086 containing one or more sulfur atoms 21/0865 {Binary compounds of nitrogen with sulfur}
21/0617	zirconium or hafnium} {with vanadium, niobium or tantalum}	_
21/0617 21/062	zirconium or hafnium} {with vanadium, niobium or tantalum} {with chromium, molybdenum or tungsten}	21/0865 {Binary compounds of nitrogen with sulfur}
21/0617	zirconium or hafnium} {with vanadium, niobium or tantalum}	21/0865 {Binary compounds of nitrogen with sulfur} 21/087 containing one or more hydrogen atoms

	(Chli : - NII Cl 1:-hli	21/20	N1:4: :
21/091	• • • {Chloramine, i.e. NH <sub>2</sub> Cl or dichloramine,	21/38	. Nitric acid
24/002	i.e. NHCl <sub>2</sub> }	21/40	• • Preparation by absorption of oxides of nitrogen
21/092	containing also one or more metal atoms	21/42	$\{(\underline{\text{C01B 21/26}} \text{ takes precedence})\}$
21/0923	{Metal imides or amides (silicon imides or	21/42	Preparation from nitrates
	amides <u>C01B 21/087</u> )}	21/44	Concentration {(C01B 21/40 takes
21/0926	{of alkali metals}	21/45	precedence)}
21/093	• • containing also one or more sulfur atoms	21/46	Purification; Separation; {Stabilisation
21/0935	• • • {Imidodisulfonic acid; Nitrilotrisulfonic		$(\underline{\text{C01B 21/40}} \text{ takes precedence})$
	acid; Salts thereof}	21/48	• • Methods for the preparation of nitrates in general
21/094	Nitrosyl containing acids		(particular individual nitrates, see the relevant
21/096	Amidosulfonic acid; Salts thereof		groups of subclasses <u>C01B</u> - <u>C01G</u> , according to
21/097	containing phosphorus atoms		the cation)
21/0975	• • {containing also one or more sulfur atoms}	21/50	Nitrous acid; Salts thereof
21/098	Phosphonitrilic dihalides; Polymers thereof	23/00	Noble gases; Compounds thereof (liquefying F25J;
21/0983	• • • {Phosphonitrilic difluorides; Polymers	20,00	{noble gases obtained by rectification <u>F25J 3/028</u> })
	thereof}	23/0005	• {Compounds of noble gases}
21/0986	• • • {Phosphonitrilic dichlorides; Polymers	23/0003	• {Purification or separation processes of noble
21/0/00	thereof}	23/001	gases}
21/12	Carbamic acid {or thiocarbamic acid}; Salts	23/0015	9 ,
21/12	thereof		• • {Chemical processing only}
21/125	• • • {Metal carbamates}	23/0021	• • · {by oxidation}
21/14	Hydroxylamine; Salts thereof	23/0026	• • · {by reduction}
21/14		23/0031	• • · {by complexation}
	· · · {Preparation}	23/0036	• • {Physical processing only}
21/1418	• • • • {by catalytic reduction of nitrogen oxides or	23/0042	• • • {by making use of membranes}
01/1/05	nitrates with hydrogen}	23/0047	• • • {characterised by the membrane}
21/1427	• • • • {by reduction of nitrogen oxides or nitrites	23/0052	• • {by adsorption in solids}
	with bisulfite or sulfur dioxide, e.g. by the	23/0057	{characterised by the adsorbent}
21/1/26	Raschig process}	23/0063	{Carbon based materials}
21/1436	• • • {by reaction in the gas phase, e.g. of	23/0068	{Zeolites}
~	nitrogen, hydrogen and oxygen}	23/0073	• • • • • {Other molecular sieve materials}
21/1445	• • • {of hydoxylamine from its salts}	23/0078	{Temperature swing adsorption}
21/1454	• • • • {of hydroxylamine salts by processes	23/0076	· · · {in getters}
	not covered by one or more of groups	23/0089	• • • {In getters} • • • {by absorption in liquids}
	<u>C01B 21/1418</u> - <u>C01B 21/1445</u> , e.g. by	23/0089	<ul><li>. {by absorption in inquits}</li><li>. {Combined chemical and physical processing}</li></ul>
			• • {Combined chemical and physical processing}
24/44/2	conversion of one salt into another}	23/0074	
21/1463	• • • {Concentration}	23/00/4	NOTE
21/1472	<ul><li> {Concentration}</li><li> {Separation}</li></ul>	23/0074	
21/1472 21/1481	<ul><li> {Concentration}</li><li> {Separation}</li><li> {Purification}</li></ul>	23/0074	In this group, processing steps are
21/1472	<ul><li> {Concentration}</li><li> {Separation}</li></ul>	23/0074	In this group, processing steps are indexed with codes chosen from
21/1472 21/1481	<ul><li> {Concentration}</li><li> {Separation}</li><li> {Purification}</li></ul>	23/0074	In this group, processing steps are
21/1472 21/1481 21/149	<ul><li> {Concentration}</li><li> {Separation}</li><li> {Purification}</li><li> {Stabilisation}</li></ul>		In this group, processing steps are indexed with codes chosen from C01B 2210/0001 - C01B 2210/0025
21/1472 21/1481 21/149 21/16	<ul> <li> {Concentration}</li> <li> {Separation}</li> <li> {Purification}</li> <li> {Stabilisation}</li> <li>. Hydrazine; Salts thereof</li> </ul>	25/00	In this group, processing steps are indexed with codes chosen from C01B 2210/0001 - C01B 2210/0025  Phosphorus; Compounds thereof ({C01B 6/00}),
21/1472 21/1481 21/149 21/16 21/20	<ul> <li> {Concentration}</li> <li> {Separation}</li> <li> {Purification}</li> <li> {Stabilisation}</li> <li>. Hydrazine; Salts thereof</li> <li>. Nitrogen oxides; Oxyacids of nitrogen; Salts thereof</li> </ul>		In this group, processing steps are indexed with codes chosen from C01B 2210/0001 - C01B 2210/0025  Phosphorus; Compounds thereof ({C01B 6/00} , C01B 21/00, C01B 23/00 take precedence;
21/1472 21/1481 21/149 21/16 21/20	<ul> <li> {Concentration}</li> <li> {Separation}</li> <li> {Purification}</li> <li> {Stabilisation}</li> <li>. Hydrazine; Salts thereof</li> <li>Nitrogen oxides; Oxyacids of nitrogen; Salts thereof</li> <li>. {Preparation of nitrogen oxides using a plasma or an electric discharge}</li> </ul>	25/00	In this group, processing steps are indexed with codes chosen from C01B 2210/0001 - C01B 2210/0025  Phosphorus; Compounds thereof ({C01B 6/00} , C01B 21/00, C01B 23/00 take precedence; perphosphates C01B 15/16)
21/1472 21/1481 21/149 21/16 21/20 21/203	<ul> <li> {Concentration}</li> <li> {Separation}</li> <li> {Purification}</li> <li> {Stabilisation}</li> <li>. Hydrazine; Salts thereof</li> <li>Nitrogen oxides; Oxyacids of nitrogen; Salts thereof</li> <li>. {Preparation of nitrogen oxides using a plasma or an electric discharge}</li> <li>. {Nitric anhydride (N<sub>2</sub>O<sub>5</sub>) (C01B 21/203 takes</li> </ul>	<b>25/00</b> 25/003	In this group, processing steps are indexed with codes chosen from C01B 2210/0001 - C01B 2210/0025  Phosphorus; Compounds thereof ({C01B 6/00} , C01B 21/00, C01B 23/00 take precedence; perphosphates C01B 15/16)  • {Phosphorus}
21/1472 21/1481 21/149 21/16 21/20 21/203 21/206	<ul> <li> {Concentration}</li> <li> {Separation}</li> <li> {Purification}</li> <li> {Stabilisation}</li> <li>. Hydrazine; Salts thereof</li> <li>Nitrogen oxides; Oxyacids of nitrogen; Salts thereof</li> <li>. {Preparation of nitrogen oxides using a plasma or an electric discharge}</li> <li>. {Nitric anhydride (N<sub>2</sub>O<sub>5</sub>) (C01B 21/203 takes precedence)}</li> </ul>	<b>25/00</b> 25/003 25/006	In this group, processing steps are indexed with codes chosen from C01B 2210/0001 - C01B 2210/0025  Phosphorus; Compounds thereof ({C01B 6/00} , C01B 21/00, C01B 23/00 take precedence; perphosphates C01B 15/16)  • {Phosphorus}  • {Stabilisation (C01B 25/04 takes precedence)}
21/1472 21/1481 21/149 21/16 21/20 21/203	<ul> <li> {Concentration}</li> <li> {Separation}</li> <li> {Purification}</li> <li> {Stabilisation}</li> <li>. Hydrazine; Salts thereof</li> <li>. Nitrogen oxides; Oxyacids of nitrogen; Salts thereof</li> <li>. {Preparation of nitrogen oxides using a plasma or an electric discharge}</li> <li>. {Nitric anhydride (N<sub>2</sub>O<sub>5</sub>) (C01B 21/203 takes precedence)}</li> <li>. Nitrous oxide (N<sub>2</sub>O) {(C01B 21/203 takes</li> </ul>	<b>25/00</b> 25/003	In this group, processing steps are indexed with codes chosen from C01B 2210/0001 - C01B 2210/0025  Phosphorus; Compounds thereof ({C01B 6/00} , C01B 21/00, C01B 23/00 take precedence; perphosphates C01B 15/16)  • {Phosphorus}  • {Stabilisation (C01B 25/04 takes precedence)}  • Treating phosphate ores or other raw phosphate
21/1472 21/1481 21/149 21/16 21/20 21/203 21/206	<ul> <li> {Concentration}</li> <li> {Separation}</li> <li> {Purification}</li> <li> {Stabilisation}</li> <li>. Hydrazine; Salts thereof</li> <li>. Nitrogen oxides; Oxyacids of nitrogen; Salts thereof</li> <li>. {Preparation of nitrogen oxides using a plasma or an electric discharge}</li> <li>. {Nitric anhydride (N<sub>2</sub>O<sub>5</sub>) (C01B 21/203 takes precedence)}</li> <li>. Nitrous oxide (N<sub>2</sub>O) {(C01B 21/203 takes precedence)}</li> </ul>	<b>25/00</b> 25/003 25/006	In this group, processing steps are indexed with codes chosen from C01B 2210/0001 - C01B 2210/0025  Phosphorus; Compounds thereof ({C01B 6/00} , C01B 21/00, C01B 23/00 take precedence; perphosphates C01B 15/16)  • {Phosphorus}  • • {Stabilisation (C01B 25/04 takes precedence)}  • Treating phosphate ores or other raw phosphate materials to obtain phosphorus or phosphorus
21/1472 21/1481 21/149 21/16 21/20 21/203 21/206	<ul> <li> {Concentration}</li> <li> {Separation}</li> <li> {Purification}</li> <li> {Stabilisation}</li> <li>. Hydrazine; Salts thereof</li> <li>. Nitrogen oxides; Oxyacids of nitrogen; Salts thereof</li> <li>. {Preparation of nitrogen oxides using a plasma or an electric discharge}</li> <li>. {Nitric anhydride (N<sub>2</sub>O<sub>5</sub>) (C01B 21/203 takes precedence)}</li> <li>. Nitrous oxide (N<sub>2</sub>O) {(C01B 21/203 takes precedence)}</li> <li>. Nitric oxide (NO) {(C01B 21/203 takes</li> </ul>	25/00 25/003 25/006 25/01	In this group, processing steps are indexed with codes chosen from C01B 2210/0001 - C01B 2210/0025  Phosphorus; Compounds thereof ({C01B 6/00} , C01B 21/00, C01B 23/00 take precedence; perphosphates C01B 15/16)  • {Phosphorus}  • • {Stabilisation (C01B 25/04 takes precedence)}  • Treating phosphate ores or other raw phosphate materials to obtain phosphorus or phosphorus compounds
21/1472 21/1481 21/149 21/16 21/20 21/203 21/206 21/22 21/24	<ul> <li> {Concentration}</li> <li> {Separation}</li> <li> {Purification}</li> <li> {Stabilisation}</li> <li>. Hydrazine; Salts thereof</li> <li>. Nitrogen oxides; Oxyacids of nitrogen; Salts thereof</li> <li>. {Preparation of nitrogen oxides using a plasma or an electric discharge}</li> <li>. {Nitric anhydride (N<sub>2</sub>O<sub>5</sub>) (C01B 21/203 takes precedence)}</li> <li>. Nitrous oxide (N<sub>2</sub>O) {(C01B 21/203 takes precedence)}</li> <li>. Nitric oxide (NO) {(C01B 21/203 takes precedence)}</li> </ul>	25/00 25/003 25/006 25/01 25/02	In this group, processing steps are indexed with codes chosen from C01B 2210/0001 - C01B 2210/0025  Phosphorus; Compounds thereof ({C01B 6/00}, C01B 21/00, C01B 23/00 take precedence; perphosphates C01B 15/16)  • {Phosphorus}  • • {Stabilisation (C01B 25/04 takes precedence)}  • Treating phosphate ores or other raw phosphate materials to obtain phosphorus or phosphorus compounds  • Preparation of phosphorus
21/1472 21/1481 21/149 21/16 21/20 21/203 21/206	<ul> <li> {Concentration}</li> <li> {Separation}</li> <li> {Purification}</li> <li> {Stabilisation}</li> <li> Hydrazine; Salts thereof</li> <li>. Nitrogen oxides; Oxyacids of nitrogen; Salts thereof</li> <li>. {Preparation of nitrogen oxides using a plasma or an electric discharge}</li> <li>. {Nitric anhydride (N<sub>2</sub>O<sub>5</sub>) (C01B 21/203 takes precedence)}</li> <li>. Nitrous oxide (N<sub>2</sub>O) {(C01B 21/203 takes precedence)}</li> <li>. Nitric oxide (NO) {(C01B 21/203 takes precedence)}</li> <li>. Nitric oxide (NO) {(C01B 21/203 takes precedence)}</li> <li> Preparation by catalytic {or non-catalytic}</li> </ul>	25/00 25/003 25/006 25/01 25/02 25/023	In this group, processing steps are indexed with codes chosen from C01B 2210/0001 - C01B 2210/0025  Phosphorus; Compounds thereof ({C01B 6/00} , C01B 21/00, C01B 23/00 take precedence; perphosphates C01B 15/16)  {Phosphorus}  • {Stabilisation (C01B 25/04 takes precedence)}  • Treating phosphate ores or other raw phosphate materials to obtain phosphorus or phosphorus compounds  • Preparation of phosphorus  • of red phosphorus
21/1472 21/1481 21/149 21/16 21/20 21/203 21/206 21/22 21/24 21/26	<ul> <li> {Concentration}</li> <li> {Separation}</li> <li> {Purification}</li> <li> {Stabilisation}</li> <li> Hydrazine; Salts thereof</li> <li>. Nitrogen oxides; Oxyacids of nitrogen; Salts thereof</li> <li>. {Preparation of nitrogen oxides using a plasma or an electric discharge}</li> <li>. {Nitric anhydride (N<sub>2</sub>O<sub>5</sub>) (C01B 21/203 takes precedence)}</li> <li>. Nitrous oxide (N<sub>2</sub>O) {(C01B 21/203 takes precedence)}</li> <li>. Nitric oxide (NO) {(C01B 21/203 takes precedence)}</li> <li>. Preparation by catalytic {or non-catalytic} oxidation of ammonia</li> </ul>	25/00 25/003 25/006 25/01 25/02	In this group, processing steps are indexed with codes chosen from C01B 2210/0001 - C01B 2210/0025  Phosphorus; Compounds thereof ({C01B 6/00} , C01B 21/00, C01B 23/00 take precedence; perphosphates C01B 15/16)  {Phosphorus}  • {Stabilisation (C01B 25/04 takes precedence)}  • Treating phosphate ores or other raw phosphate materials to obtain phosphorus or phosphorus compounds  • Preparation of phosphorus  • of red phosphorus  • of yellow phosphorus
21/1472 21/1481 21/149 21/16 21/20 21/203 21/206 21/22 21/24 21/26 21/26	<ul> <li> {Concentration}</li> <li> {Separation}</li> <li> {Purification}</li> <li> {Stabilisation}</li> <li> Hydrazine; Salts thereof</li> <li>. Nitrogen oxides; Oxyacids of nitrogen; Salts thereof</li> <li>. {Preparation of nitrogen oxides using a plasma or an electric discharge}</li> <li>. {Nitric anhydride (N<sub>2</sub>O<sub>5</sub>) (C01B 21/203 takes precedence)}</li> <li>. Nitrous oxide (N<sub>2</sub>O) {(C01B 21/203 takes precedence)}</li> <li>. Nitric oxide (NO) {(C01B 21/203 takes precedence)}</li> <li>. Nitric oxide (NO) {(C01B 21/203 takes precedence)}</li> <li> Preparation by catalytic {or non-catalytic} oxidation of ammonia</li> <li> {obtaining nitrogen dioxide or tetroxide}</li> </ul>	25/00 25/003 25/006 25/01 25/02 25/023	In this group, processing steps are indexed with codes chosen from C01B 2210/0001 - C01B 2210/0025  Phosphorus; Compounds thereof ({C01B 6/00} , C01B 21/00, C01B 23/00 take precedence; perphosphates C01B 15/16)  {Phosphorus}  • {Stabilisation (C01B 25/04 takes precedence)}  • Treating phosphate ores or other raw phosphate materials to obtain phosphorus or phosphorus compounds  • Preparation of phosphorus  • of red phosphorus
21/1472 21/1481 21/149 21/16 21/20 21/203 21/206 21/22 21/24 21/26 21/262 21/265	<ul> <li> {Concentration}</li> <li> {Separation}</li> <li> {Purification}</li> <li> {Stabilisation}</li> <li>. Hydrazine; Salts thereof</li> <li>. Nitrogen oxides; Oxyacids of nitrogen; Salts thereof</li> <li>. {Preparation of nitrogen oxides using a plasma or an electric discharge}</li> <li>. {Nitric anhydride (N<sub>2</sub>O<sub>5</sub>) (C01B 21/203 takes precedence)}</li> <li>. Nitrous oxide (N<sub>2</sub>O) {(C01B 21/203 takes precedence)}</li> <li>. Nitric oxide (NO) {(C01B 21/203 takes precedence)}</li> <li>. Nitric oxide (NO) {(C01B 21/203 takes precedence)}</li> <li> Preparation by catalytic {or non-catalytic} oxidation of ammonia</li> <li> {obtaining nitrogen dioxide or tetroxide}</li> <li> {characterised by the catalyst}</li> </ul>	25/00 25/003 25/006 25/01 25/02 25/023 25/027	In this group, processing steps are indexed with codes chosen from C01B 2210/0001 - C01B 2210/0025  Phosphorus; Compounds thereof ({C01B 6/00} , C01B 21/00, C01B 23/00 take precedence; perphosphates C01B 15/16)  {Phosphorus}  • {Stabilisation (C01B 25/04 takes precedence)}  • Treating phosphate ores or other raw phosphate materials to obtain phosphorus or phosphorus compounds  • Preparation of phosphorus  • of red phosphorus  • of yellow phosphorus
21/1472 21/1481 21/149 21/16 21/20 21/203 21/206 21/22 21/24 21/26 21/26	<ul> <li> {Concentration}</li> <li> {Separation}</li> <li> {Purification}</li> <li> {Stabilisation}</li> <li> Hydrazine; Salts thereof</li> <li>. Nitrogen oxides; Oxyacids of nitrogen; Salts thereof</li> <li>. {Preparation of nitrogen oxides using a plasma or an electric discharge}</li> <li>. {Nitric anhydride (N<sub>2</sub>O<sub>5</sub>) (C01B 21/203 takes precedence)}</li> <li>. Nitrous oxide (N<sub>2</sub>O) {(C01B 21/203 takes precedence)}</li> <li>. Nitric oxide (NO) {(C01B 21/203 takes precedence)}</li> <li>. Preparation by catalytic {or non-catalytic} oxidation of ammonia</li> <li> {obtaining nitrogen dioxide or tetroxide}</li> <li> {characterised by the catalyst}</li> <li> {Means for preventing deterioration or loss</li> </ul>	25/00 25/003 25/006 25/01 25/02 25/023 25/027 25/04	In this group, processing steps are indexed with codes chosen from C01B 2210/0001 - C01B 2210/0025  Phosphorus; Compounds thereof ({C01B 6/00}, C01B 21/00, C01B 23/00 take precedence; perphosphates C01B 15/16)  • {Phosphorus}  • • {Stabilisation (C01B 25/04 takes precedence)}  • Treating phosphate ores or other raw phosphate materials to obtain phosphorus or phosphorus compounds  • Preparation of phosphorus  • • of red phosphorus  • of yellow phosphorus  • Purification of phosphorus
21/1472 21/1481 21/149 21/16 21/20 21/203 21/206 21/22 21/24 21/26 21/262 21/265 21/267	<ul> <li> {Concentration}</li> <li> {Separation}</li> <li> {Purification}</li> <li> {Stabilisation}</li> <li>. Hydrazine; Salts thereof</li> <li>. Nitrogen oxides; Oxyacids of nitrogen; Salts thereof</li> <li>. {Preparation of nitrogen oxides using a plasma or an electric discharge}</li> <li>. {Nitric anhydride (N<sub>2</sub>O<sub>5</sub>) (C01B 21/203 takes precedence)}</li> <li>. Nitrous oxide (N<sub>2</sub>O) {(C01B 21/203 takes precedence)}</li> <li>. Nitric oxide (NO) {(C01B 21/203 takes precedence)}</li> <li>. Nitric oxide (NO) {(C01B 21/203 takes precedence)}</li> <li> Preparation by catalytic {or non-catalytic} oxidation of ammonia</li> <li> {obtaining nitrogen dioxide or tetroxide}</li> <li> {characterised by the catalyst}</li> <li> {Means for preventing deterioration or loss of catalyst or for recovering lost catalyst}</li> </ul>	25/00 25/003 25/006 25/01 25/02 25/023 25/027 25/04 25/043	In this group, processing steps are indexed with codes chosen from C01B 2210/0001 - C01B 2210/0025  Phosphorus; Compounds thereof ({C01B 6/00} , C01B 21/00, C01B 23/00 take precedence; perphosphates C01B 15/16)  • {Phosphorus}  • • {Stabilisation (C01B 25/04 takes precedence)}  • Treating phosphate ores or other raw phosphate materials to obtain phosphorus or phosphorus compounds  • Preparation of phosphorus  • • of red phosphorus  • of yellow phosphorus  • Purification of phosphorus  • of red phosphorus
21/1472 21/1481 21/149 21/16 21/20 21/203 21/206 21/22 21/24 21/26 21/262 21/265 21/267	<ul> <li> {Concentration}</li> <li> {Separation}</li> <li> {Purification}</li> <li> {Stabilisation}</li> <li> Hydrazine; Salts thereof</li> <li>. Nitrogen oxides; Oxyacids of nitrogen; Salts thereof</li> <li>. {Preparation of nitrogen oxides using a plasma or an electric discharge}</li> <li>. {Nitric anhydride (N<sub>2</sub>O<sub>5</sub>) (C01B 21/203 takes precedence)}</li> <li>. Nitrous oxide (N<sub>2</sub>O) {(C01B 21/203 takes precedence)}</li> <li>. Nitric oxide (NO) {(C01B 21/203 takes precedence)}</li> <li>. Nitric oxide (NO) {(C01B 21/203 takes precedence)}</li> <li> Preparation by catalytic {or non-catalytic} oxidation of ammonia</li> <li> {obtaining nitrogen dioxide or tetroxide}</li> <li> {characterised by the catalyst}</li> <li> {Means for preventing deterioration or loss of catalyst or for recovering lost catalyst}</li> <li> Apparatus</li> </ul>	25/00 25/003 25/006 25/01 25/02 25/023 25/027 25/04 25/043 25/047	In this group, processing steps are indexed with codes chosen from C01B 2210/0001 - C01B 2210/0025  Phosphorus; Compounds thereof ({C01B 6/00} , C01B 21/00, C01B 23/00 take precedence; perphosphates C01B 15/16)  • {Phosphorus}  • • {Stabilisation (C01B 25/04 takes precedence)}  • Treating phosphate ores or other raw phosphate materials to obtain phosphorus or phosphorus compounds  • Preparation of phosphorus  • • of red phosphorus  • of yellow phosphorus  • of red phosphorus  • of red phosphorus  • of yellow phosphorus  • of yellow phosphorus  • of yellow phosphorus  • Hydrogen phosphides
21/1472 21/1481 21/149 21/16 21/20 21/203 21/206 21/22 21/24 21/26 21/262 21/265 21/267	<ul> <li> {Concentration}</li> <li> {Separation}</li> <li> {Purification}</li> <li> {Stabilisation}</li> <li> Hydrazine; Salts thereof</li> <li>. Nitrogen oxides; Oxyacids of nitrogen; Salts thereof</li> <li>. {Preparation of nitrogen oxides using a plasma or an electric discharge}</li> <li>. {Nitric anhydride (N<sub>2</sub>O<sub>5</sub>) (C01B 21/203 takes precedence)}</li> <li>. Nitrous oxide (N<sub>2</sub>O) {(C01B 21/203 takes precedence)}</li> <li>. Nitric oxide (NO) {(C01B 21/203 takes precedence)}</li> <li>. Preparation by catalytic {or non-catalytic} oxidation of ammonia</li> <li> {obtaining nitrogen dioxide or tetroxide}</li> <li> {characterised by the catalyst}</li> <li> {Means for preventing deterioration or loss of catalyst or for recovering lost catalyst}</li> <li> Apparatus</li> <li>. Preparation by oxidation of nitrogen</li> </ul>	25/00 25/003 25/006 25/01 25/02 25/023 25/027 25/04 25/043 25/047 25/06 25/08	In this group, processing steps are indexed with codes chosen from C01B 2210/0001 - C01B 2210/0025  Phosphorus; Compounds thereof ({C01B 6/00}, C01B 21/00, C01B 23/00 take precedence; perphosphates C01B 15/16)  • {Phosphorus}  • • {Stabilisation (C01B 25/04 takes precedence)}  • Treating phosphate ores or other raw phosphate materials to obtain phosphorus or phosphorus compounds  • Preparation of phosphorus  • • of red phosphorus  • • of yellow phosphorus  • • Hydrogen phosphides  • Other phosphides
21/1472 21/1481 21/149 21/16 21/20 21/203 21/206 21/22 21/24 21/26 21/262 21/265 21/267 21/28 21/30	<ul> <li> {Concentration}</li> <li> {Separation}</li> <li> {Purification}</li> <li> {Stabilisation}</li> <li> Hydrazine; Salts thereof</li> <li>. Nitrogen oxides; Oxyacids of nitrogen; Salts thereof</li> <li>. {Preparation of nitrogen oxides using a plasma or an electric discharge}</li> <li>. {Nitric anhydride (N<sub>2</sub>O<sub>5</sub>) (C01B 21/203 takes precedence)}</li> <li>. Nitrous oxide (N<sub>2</sub>O) {(C01B 21/203 takes precedence)}</li> <li>. Nitric oxide (NO) {(C01B 21/203 takes precedence)}</li> <li>. Preparation by catalytic {or non-catalytic} oxidation of ammonia</li> <li> {obtaining nitrogen dioxide or tetroxide}</li> <li> {Characterised by the catalyst}</li> <li> {Means for preventing deterioration or loss of catalyst or for recovering lost catalyst}</li> <li> Apparatus</li> <li> Preparation by oxidation of nitrogen {(C01B 21/26 takes precedence)}</li> </ul>	25/00 25/003 25/006 25/01 25/02 25/023 25/027 25/04 25/043 25/047 25/06	In this group, processing steps are indexed with codes chosen from C01B 2210/0001 - C01B 2210/0025  Phosphorus; Compounds thereof ({C01B 6/00}, C01B 21/00, C01B 23/00 take precedence; perphosphates C01B 15/16)  • {Phosphorus}  • • {Stabilisation (C01B 25/04 takes precedence)}  • Treating phosphate ores or other raw phosphate materials to obtain phosphorus or phosphorus compounds  • Preparation of phosphorus  • • of red phosphorus  • • of yellow phosphorus  • • of phosphorus  • • of yellow phosphorus
21/1472 21/1481 21/149 21/16 21/20 21/203 21/206 21/22 21/24 21/26 21/262 21/265 21/267 21/28 21/30	<ul> <li> {Concentration}</li> <li> {Separation}</li> <li> {Purification}</li> <li> {Stabilisation}</li> <li> Hydrazine; Salts thereof</li> <li>. Nitrogen oxides; Oxyacids of nitrogen; Salts thereof</li> <li>. {Preparation of nitrogen oxides using a plasma or an electric discharge}</li> <li>. {Nitric anhydride (N<sub>2</sub>O<sub>5</sub>) (C01B 21/203 takes precedence)}</li> <li>. Nitrous oxide (N<sub>2</sub>O) {(C01B 21/203 takes precedence)}</li> <li>. Nitric oxide (NO) {(C01B 21/203 takes precedence)}</li> <li>. Preparation by catalytic {or non-catalytic} oxidation of ammonia</li> <li> {obtaining nitrogen dioxide or tetroxide}</li> <li> {Means for preventing deterioration or loss of catalyst or for recovering lost catalyst}</li> <li> Apparatus</li> <li> Preparation by oxidation of nitrogen {(C01B 21/26 takes precedence)}</li> <li> Apparatus</li> <li> Apparatus</li> </ul>	25/00 25/003 25/006 25/01 25/02 25/023 25/027 25/04 25/043 25/047 25/06 25/08 25/081	In this group, processing steps are indexed with codes chosen from C01B 2210/0001 - C01B 2210/0025  Phosphorus; Compounds thereof ({C01B 6/00}, C01B 21/00, C01B 23/00 take precedence; perphosphates C01B 15/16)  • {Phosphorus}  • • {Stabilisation (C01B 25/04 takes precedence)}  • Treating phosphate ores or other raw phosphate materials to obtain phosphorus or phosphorus compounds  • Preparation of phosphorus  • • of red phosphorus  • • of yellow phosphorus  • • of jellow phosphorus  • • of yellow phosphorus  • • of jellow phosphorus  • • of jellow phosphides  • • (of alkali metals, alkaline-earth metals or magnesium)
21/1472 21/1481 21/149 21/16 21/20 21/203 21/206 21/22 21/24 21/26 21/262 21/265 21/267 21/28 21/30	<ul> <li> {Concentration}</li> <li> {Separation}</li> <li> {Purification}</li> <li> {Stabilisation}</li> <li> {Stabilisation}</li> <li> Hydrazine; Salts thereof</li> <li>. Nitrogen oxides; Oxyacids of nitrogen; Salts thereof</li> <li>. {Preparation of nitrogen oxides using a plasma or an electric discharge}</li> <li>. {Nitric anhydride (N<sub>2</sub>O<sub>5</sub>) (C01B 21/203 takes precedence)}</li> <li>. Nitrous oxide (N<sub>2</sub>O) {(C01B 21/203 takes precedence)}</li> <li>. Nitric oxide (NO) {(C01B 21/203 takes precedence)}</li> <li> Preparation by catalytic {or non-catalytic} oxidation of ammonia</li> <li> {obtaining nitrogen dioxide or tetroxide}</li> <li> {Characterised by the catalyst}</li> <li> {Means for preventing deterioration or loss of catalyst or for recovering lost catalyst}</li> <li> Apparatus</li> <li> Preparation by oxidation of nitrogen {(C01B 21/26 takes precedence)}</li> <li> Apparatus</li> <li>. Nitrogen trioxide (N<sub>2</sub>O<sub>3</sub>) {(C01B 21/203 takes</li> </ul>	25/00 25/003 25/006 25/01 25/02 25/023 25/027 25/04 25/043 25/047 25/06 25/08 25/081	In this group, processing steps are indexed with codes chosen from C01B 2210/0001 - C01B 2210/0025  Phosphorus; Compounds thereof ({C01B 6/00}, C01B 21/00, C01B 23/00 take precedence; perphosphates C01B 15/16)  • {Phosphorus}  • • {Stabilisation (C01B 25/04 takes precedence)}  • Treating phosphate ores or other raw phosphate materials to obtain phosphorus or phosphorus compounds  • Preparation of phosphorus  • • of red phosphorus  • of yellow phosphorus  • of of alkali metals, alkaline-earth metals or magnesium}  • • {of boron, aluminium, gallium or indium}
21/1472 21/1481 21/149 21/16 21/20 21/203 21/206 21/22 21/24 21/26 21/265 21/265 21/267 21/28 21/30 21/32 21/34	<ul> <li>. (Concentration)</li> <li>. (Separation)</li> <li>. (Purification)</li> <li>. (Stabilisation)</li> <li>. Hydrazine; Salts thereof</li> <li>. Nitrogen oxides; Oxyacids of nitrogen; Salts thereof</li> <li>. (Preparation of nitrogen oxides using a plasma or an electric discharge)</li> <li>. (Nitric anhydride (N<sub>2</sub>O<sub>5</sub>) (C01B 21/203 takes precedence))</li> <li>. Nitrous oxide (N<sub>2</sub>O) {(C01B 21/203 takes precedence)}</li> <li>. Nitric oxide (NO) {(C01B 21/203 takes precedence)}</li> <li>. Preparation by catalytic {or non-catalytic} oxidation of ammonia</li> <li>. (obtaining nitrogen dioxide or tetroxide)</li> <li>. (characterised by the catalyst)</li> <li>. (Means for preventing deterioration or loss of catalyst or for recovering lost catalyst)</li> <li>. Apparatus</li> <li>. Preparation by oxidation of nitrogen {(C01B 21/26 takes precedence)}</li> <li>. Apparatus</li> <li>. Nitrogen trioxide (N<sub>2</sub>O<sub>3</sub>) {(C01B 21/203 takes precedence)}</li> <li>. Apparatus</li> <li>. Nitrogen trioxide (N<sub>2</sub>O<sub>3</sub>) {(C01B 21/203 takes precedence)}</li> </ul>	25/00 25/003 25/006 25/01 25/02 25/023 25/027 25/04 25/043 25/047 25/06 25/08 25/081 25/082 25/084	In this group, processing steps are indexed with codes chosen from C01B 2210/0001 - C01B 2210/0025  Phosphorus; Compounds thereof ({C01B 6/00}, C01B 21/00, C01B 23/00 take precedence; perphosphates C01B 15/16)  • {Phosphorus}  • • {Stabilisation (C01B 25/04 takes precedence)}  • Treating phosphate ores or other raw phosphate materials to obtain phosphorus or phosphorus compounds  • Preparation of phosphorus  • • of red phosphorus  • of yellow phosphorus  • of yellow phosphorus  • of red phosphorus  • of yellow phosphorus  • of yellow phosphorus  • of of alkali metals, alkaline-earth metals or magnesium}  • • {of boron, aluminium, gallium or indium}  • • • {of boron}
21/1472 21/1481 21/149 21/16 21/20 21/203 21/206 21/22 21/24 21/26 21/262 21/265 21/267 21/28 21/30	<ul> <li> {Concentration}</li> <li> {Separation}</li> <li> {Purification}</li> <li> {Stabilisation}</li> <li> {Stabilisation}</li> <li> Hydrazine; Salts thereof</li> <li>. Nitrogen oxides; Oxyacids of nitrogen; Salts thereof</li> <li>. {Preparation of nitrogen oxides using a plasma or an electric discharge}</li> <li>. {Nitric anhydride (N<sub>2</sub>O<sub>5</sub>) (C01B 21/203 takes precedence)}</li> <li>. Nitrous oxide (N<sub>2</sub>O) {(C01B 21/203 takes precedence)}</li> <li>. Nitric oxide (NO) {(C01B 21/203 takes precedence)}</li> <li> Preparation by catalytic {or non-catalytic} oxidation of ammonia</li> <li> {obtaining nitrogen dioxide or tetroxide}</li> <li> {Characterised by the catalyst}</li> <li> {Means for preventing deterioration or loss of catalyst or for recovering lost catalyst}</li> <li> Apparatus</li> <li> Preparation by oxidation of nitrogen {(C01B 21/26 takes precedence)}</li> <li> Apparatus</li> <li>. Nitrogen trioxide (N<sub>2</sub>O<sub>3</sub>) {(C01B 21/203 takes</li> </ul>	25/00 25/003 25/006 25/01 25/02 25/023 25/027 25/04 25/043 25/047 25/06 25/08 25/081	In this group, processing steps are indexed with codes chosen from C01B 2210/0001 - C01B 2210/0025  Phosphorus; Compounds thereof ({C01B 6/00}, C01B 21/00, C01B 23/00 take precedence; perphosphates C01B 15/16)  • {Phosphorus}  • • {Stabilisation (C01B 25/04 takes precedence)}  • Treating phosphate ores or other raw phosphate materials to obtain phosphorus or phosphorus compounds  • Preparation of phosphorus  • • of red phosphorus  • of yellow phosphorus  • of of alkali metals, alkaline-earth metals or magnesium}  • • {of boron, aluminium, gallium or indium}

25/088	• • {containing plural metal}	25/235 Clarification; Stabilisation to prevent post-
25/10	Halides or oxyhalides of phosphorus	precipitation of dissolved impurities
25/12	• Oxides of phosphorus	25/237 Selective elimination of impurities
25/14	Sulfur, selenium, or tellurium compounds of	$\{(\underbrace{\text{C01B 25/2343}}_{\text{25/2372}} \text{ takes precedence})\}$
25/16	phosphorus	25/2372 { Anionic impurities, e.g. silica or boron compounds }
25/16	• Oxyacids of phosphorus; Salts thereof (peroxyacids	25/2375 {Fluoride or fluosilicate anion}
05/161	or salts thereof <u>C01B 15/00</u> )	
25/161	<ul> <li>{containing at least one phosphorus atom with an oxidation number less than five, other than those</li> </ul>	,
	mentioned below; Salts thereof}	25/238 Cationic impurities, {e.g. arsenic compounds}
25/163	Phosphorous acid; Salts thereof	25/24 Condensed phosphoric acids
25/165	Hypophosphorous acid; Salts thereof	25/26 • Phosphates (perphosphates C01B 15/16)
25/168	<ul> <li>Pyrophosphorous acid; Salts thereof</li> <li>Pyrophosphorous acid; Salts thereof</li> </ul>	25/265 • • • General methods for obtaining phosphates
25/18	Phosphoric acid	25/28 Ammonium phosphates
25/185	Preparation neither from elemental	25/30 Alkali metal phosphates
23/103	phosphorus or phosphoric anhydride nor by	25/301 {Preparation from liquid orthophosphoric
	reacting phosphate-containing material with	acid or from an acid solution or suspension
	an acid, e.g. by reacting phosphate-containing	of orthophosphates (using ion-exchangers
	material with an ion-exchange resin or an acid	C01B 25/30)}
	salt used alone}	25/303 {with elimination of impurities}
25/20	Preparation from elemental phosphorus or	25/305 {Preparation from phosphorus-containing
	phosphoric anhydride	compounds by alkaline treatment}
25/22	• • • Preparation by reacting phosphate-containing	25/306 {from phosphates}
	material with an acid, e.g. wet process	25/308 {Methods for converting an alkali
25/2204	• • • • {Arrangements of vessels used in reacting	metal orthophosphate into another one;
	phosphate-containing material in wet	Purification; Decolorasing; Dehydrating;
	process}	Drying}
25/2208	• • • { with an acid or a mixture of acids other than	25/32 Phosphates of magnesium, calcium, strontium,
25/2212	sulfuric acid}	or barium
25/2212	• • • • {with hydrochloric acid or hydrogen	25/321 {Methods for converting an alkaline earth
25/2216	chloride in aqueous medium}	metal ortho-phosphate into another ortho-
25/2216	• • • • { with nitric acid or nitrous vapours in aqueous medium }	phosphate (by reaction, e.g. of phosphate
25/222	• • • with sulfuric acid, a mixture of acids mainly	rock with phosphoric acid C01B 25/322)}
23/222	consisting of sulfuric acid or a mixture of	25/322 {Preparation by neutralisation of orthophosphoric acid}
	compounds forming it <u>in situ</u> , e.g. a mixture	
	of sulfur dioxide, water and oxygen	25/324 {Preparation from a reaction solution obtained by acidifying with an acid other
25/223	only one form of calcium sulfate being	than orthophosphoric acid}
	formed	25/325 {Preparation by double decomposition}
25/2235	{Anhydrite processes}	25/327 {After-treatment (increasing the phosphate
25/225	Dihydrate process	content of ores C01B 25/32)}
25/226	Hemihydrate process	25/328 {Defluorination during or after the
25/228	one form of calcium sulfate being formed	preparation}
	and then converted to another form	25/34 Magnesium phosphates
25/2285	{Dihydrate-anhydrite or hemihydrate-	25/36 Aluminium phosphates
	anhydrite process}	25/37 Phosphates of heavy metals
25/229	Hemihydrate-dihydrate process	25/372 {of titanium, vanadium, zirconium, niobium,
25/2295	• • • • • { the conversion being performed	hafnium or tantalum}
	in one or more vessels different	25/375 {of iron}
	from those used for reaction after	25/377 { of manganese }
	separation of phosphoric acid}	25/38 Condensed phosphates
25/231	Dihydrate-hemihydrate process	25/385 {of alkaline-earth metals or magnesium}
25/232	Preparation by reacting phosphate	25/39 of alkali metals
	containing material with concentrated	25/395 {Preparation and dehydrating}
	sulfuric acid and subsequently lixiviating	25/40 Polyphosphates
25/234	the obtained mass, e.g. clinker process  • • • Purification; Stabilisation; Concentration	25/405 {of ammonium}
<i>23/2</i> 34	(purification concomitant with preparation	25/41 of alkali metals
	<u>C01B 25/22;</u> preparation involving solvent-	25/412 {Preparation from alkali metal
	solvent extraction C01B 25/46)	orthophosphates}
25/2343	{Concentration concomitant with	25/414 {Apparatus}
25, 25 75	purification, e.g. elimination of fluorine}	25/416 {Pure alkali metal polyphosphates
25/2346	{Concentration}	from impure starting materials}
		25/418 {After-treatment}

05/40	D 1 1	21/005	G 1 (GE) (GE)
25/42	Pyrophosphates		Carbon fluorides, e.g. (CF)n or (C2F)n
25/425	• • • • {of alkali metals}		reparation of intercalation compounds of graphite
25/44	Metaphosphates		ith fluorine <u>C01B 31/0415</u> )}
25/445	of alkali metals		reparation of carbon (by using ultra high pressure,
25/45	containing plural metal, or metal and		g. for the formation of diamonds, <u>B01J 3/06</u> ;
	ammonium		crystal growth <u>C30B</u> ); Purification; {After-
25/451	• • • {containing metal and ammonium}		eatment}
25/453	• • • • {having molecular-sieve properties}	31/0206	{Nanosized carbon materials (graphene
	<u>WARNING</u>	21/0212	<u>C01B 31/0438</u> )}
	WARRING		• {Fullerenes}
	Group C01B 25/453 is no longer used for	31/022	• {Carbon nanotubes}
	the classification of new documents from		NOTE
	May, 1995. The backlog of this groups		
	is continuously being reclassified to the		In groups <u>C01B 31/022</u> - <u>C01B 31/0293</u>
	appropriate subgroups of C01B 37/00 and		it is desirable to add indexing codes for
	<u>C01B 39/00</u> .		structural aspects or properties of carbon
25/455	• • • containing halogen {(completely halogenated		nanotubes. The indexing codes are chosen
23/433	alkali metal phosphates <u>C01D</u> , e.g. lithium		from <u>C01B 2202/00</u> - <u>C01B 2202/36</u>
	hexafluorophosphate C01D 15/005)}	31/0226	• • {Preparation}
25/4555	• • • • {Hypochlorite-phosphate double salts, e.g.	31/0233	• • {characterized by the catalyst}
207 .000	4(Na3PO411H2O). NaOCl or so-called		• • • {being a continuous process}
	chlorinated trisodium phosphate}		• • {in the liquid phase}
25/46	Preparation involving solvent-solvent extraction		• • {After-treatments}
	(solvent extraction in general <u>B01D 11/00</u> )		• • {Purification}
25/461	• • • {the phosphoric acid present in the medium		• • {Sorting}
	obtained after reaction being first extracted		• • • {Derivatisation, solubilisation or
	from the liquid phase formed or separated	31/02/3	dispersion in solvents}
	then re-extracted as free acid by using water	31/028	{Cutting}
	or as a phosphate by using a basic compound		{Opening or filling}
	(selective extraction of impurities contained in		• {Other structures, e.g. nano-onions, nano-
	acid <u>C01B 25/237</u> )}	31/02/3	scrolls, nano-horns, nano-cones or nano-walls)
			serons, nano nome, nano conce or nano wans,
	NOTES	31/04	Graphite including modified graphite e.g.
	<u>NOTES</u>	31/04	Graphite, including modified graphite, e.g.
	The extracting agent may be diluted with	31/04	graphitic oxides, intercalated graphite, expanded
	The extracting agent may be diluted with a compound or a mixture of compounds		graphitic oxides, intercalated graphite, expanded graphite or graphene
	1. The extracting agent may be diluted with a compound or a mixture of compounds which are not solvents for phosphoric acid,		graphitic oxides, intercalated graphite, expanded
	The extracting agent may be diluted with a compound or a mixture of compounds		graphitic oxides, intercalated graphite, expanded graphite or graphene  NOTE
	1. The extracting agent may be diluted with a compound or a mixture of compounds which are not solvents for phosphoric acid,		graphitic oxides, intercalated graphite, expanded graphite or graphene  NOTE  In groups C01B 31/04 - C01B 31/0492
	1. The extracting agent may be diluted with a compound or a mixture of compounds which are not solvents for phosphoric acid, e.g. a hydrocarbon		graphitic oxides, intercalated graphite, expanded graphite or graphene  NOTE
	<ol> <li>The extracting agent may be diluted with a compound or a mixture of compounds which are not solvents for phosphoric acid, e.g. a hydrocarbon</li> <li>Documents which belong to more than one</li> </ol>		graphitic oxides, intercalated graphite, expanded graphite or graphene  NOTE  In groups C01B 31/04 - C01B 31/0492 it is desirable to add indexing codes for
	<ol> <li>The extracting agent may be diluted with a compound or a mixture of compounds which are not solvents for phosphoric acid, e.g. a hydrocarbon</li> <li>Documents which belong to more than one subgroup of C01B 25/462 - C01B 25/466</li> </ol>		graphitic oxides, intercalated graphite, expanded graphite or graphene  NOTE  In groups C01B 31/04 - C01B 31/0492 it is desirable to add indexing codes for structural aspects or properties of graphene.
25/462	<ol> <li>The extracting agent may be diluted with a compound or a mixture of compounds which are not solvents for phosphoric acid, e.g. a hydrocarbon</li> <li>Documents which belong to more than one subgroup of C01B 25/462 - C01B 25/466 are classified by a combination, e.g. C01B 25/462 +B4+B8</li> </ol>		graphitic oxides, intercalated graphite, expanded graphite or graphene  NOTE  In groups C01B 31/04 - C01B 31/0492 it is desirable to add indexing codes for structural aspects or properties of graphene. The indexing codes are chosen from C01B 2204/00 - C01B 2204/32
25/462	<ol> <li>The extracting agent may be diluted with a compound or a mixture of compounds which are not solvents for phosphoric acid, e.g. a hydrocarbon</li> <li>Documents which belong to more than one subgroup of C01B 25/462 - C01B 25/466 are classified by a combination, e.g. C01B 25/462 +B4+B8</li> <li> {the extracting agent being alcohol or a</li> </ol>		graphitic oxides, intercalated graphite, expanded graphite or graphene  NOTE  In groups C01B 31/04 - C01B 31/0492 it is desirable to add indexing codes for structural aspects or properties of graphene. The indexing codes are chosen from C01B 2204/00 - C01B 2204/32  • {Purification; Recovery or purification of
	<ol> <li>The extracting agent may be diluted with a compound or a mixture of compounds which are not solvents for phosphoric acid, e.g. a hydrocarbon</li> <li>Documents which belong to more than one subgroup of C01B 25/462 - C01B 25/466 are classified by a combination, e.g. C01B 25/462 +B4+B8</li> <li> {the extracting agent being alcohol or a mixture of alcohols}</li> </ol>		graphitic oxides, intercalated graphite, expanded graphite or graphene  NOTE  In groups C01B 31/04 - C01B 31/0492 it is desirable to add indexing codes for structural aspects or properties of graphene. The indexing codes are chosen from C01B 2204/00 - C01B 2204/32  • {Purification; Recovery or purification of graphite formed in iron making, e.g. kish
25/462 25/463	<ol> <li>The extracting agent may be diluted with a compound or a mixture of compounds which are not solvents for phosphoric acid, e.g. a hydrocarbon</li> <li>Documents which belong to more than one subgroup of C01B 25/462 - C01B 25/466 are classified by a combination, e.g. C01B 25/462 +B4+B8</li> <li> {the extracting agent being alcohol or a mixture of alcohols}</li> <li> {the extracting agent being a ketone or a</li> </ol>	31/0407	graphitic oxides, intercalated graphite, expanded graphite or graphene  NOTE  In groups C01B 31/04 - C01B 31/0492 it is desirable to add indexing codes for structural aspects or properties of graphene. The indexing codes are chosen from C01B 2204/00 - C01B 2204/32  • {Purification; Recovery or purification of graphite formed in iron making, e.g. kish graphite}
25/463	<ol> <li>The extracting agent may be diluted with a compound or a mixture of compounds which are not solvents for phosphoric acid, e.g. a hydrocarbon</li> <li>Documents which belong to more than one subgroup of C01B 25/462 - C01B 25/466 are classified by a combination, e.g. C01B 25/462 +B4+B8</li> <li> {the extracting agent being alcohol or a mixture of alcohols}</li> <li> {the extracting agent being a ketone or a mixture of ketones}</li> </ol>	31/0407	graphitic oxides, intercalated graphite, expanded graphite or graphene  NOTE  In groups C01B 31/04 - C01B 31/0492 it is desirable to add indexing codes for structural aspects or properties of graphene. The indexing codes are chosen from C01B 2204/00 - C01B 2204/32  • {Purification; Recovery or purification of graphite formed in iron making, e.g. kish graphite}  • {Intercalation}
	<ol> <li>The extracting agent may be diluted with a compound or a mixture of compounds which are not solvents for phosphoric acid, e.g. a hydrocarbon</li> <li>Documents which belong to more than one subgroup of C01B 25/462 - C01B 25/466 are classified by a combination, e.g. C01B 25/462 +B4+B8</li> <li> {the extracting agent being alcohol or a mixture of alcohols}</li> <li> {the extracting agent being a ketone or a mixture of ketones}</li> <li> {the extracting agent being an ether or a</li> </ol>	31/0407	graphitic oxides, intercalated graphite, expanded graphite or graphene  NOTE  In groups C01B 31/04 - C01B 31/0492 it is desirable to add indexing codes for structural aspects or properties of graphene. The indexing codes are chosen from C01B 2204/00 - C01B 2204/32  • {Purification; Recovery or purification of graphite formed in iron making, e.g. kish graphite}
25/463 25/464	<ol> <li>The extracting agent may be diluted with a compound or a mixture of compounds which are not solvents for phosphoric acid, e.g. a hydrocarbon</li> <li>Documents which belong to more than one subgroup of C01B 25/462 - C01B 25/466 are classified by a combination, e.g. C01B 25/462 +B4+B8</li> <li> {the extracting agent being alcohol or a mixture of alcohols}</li> <li> {the extracting agent being a ketone or a mixture of ketones}</li> <li> {the extracting agent being an ether or a mixture of ethers}</li> </ol>	31/0407	graphitic oxides, intercalated graphite, expanded graphite or graphene  NOTE  In groups C01B 31/04 - C01B 31/0492 it is desirable to add indexing codes for structural aspects or properties of graphene. The indexing codes are chosen from C01B 2204/00 - C01B 2204/32  • {Purification; Recovery or purification of graphite formed in iron making, e.g. kish graphite}  • {Intercalation}  WARNING
25/463	<ol> <li>The extracting agent may be diluted with a compound or a mixture of compounds which are not solvents for phosphoric acid, e.g. a hydrocarbon</li> <li>Documents which belong to more than one subgroup of C01B 25/462 - C01B 25/466 are classified by a combination, e.g. C01B 25/462 +B4+B8</li> <li>(the extracting agent being alcohol or a mixture of alcohols)</li> <li>(the extracting agent being a ketone or a mixture of ketones)</li> <li>(the extracting agent being an ether or a mixture of ethers)</li> <li>(the extracting agent being an ester or a</li> </ol>	31/0407	graphitic oxides, intercalated graphite, expanded graphite or graphene  NOTE  In groups C01B 31/04 - C01B 31/0492 it is desirable to add indexing codes for structural aspects or properties of graphene. The indexing codes are chosen from C01B 2204/00 - C01B 2204/32  • {Purification; Recovery or purification of graphite formed in iron making, e.g. kish graphite}  • {Intercalation}  WARNING  Group C01B 31/0415 is not complete
25/463 25/464 25/465	<ol> <li>The extracting agent may be diluted with a compound or a mixture of compounds which are not solvents for phosphoric acid, e.g. a hydrocarbon</li> <li>Documents which belong to more than one subgroup of C01B 25/462 - C01B 25/466 are classified by a combination, e.g. C01B 25/462 +B4+B8</li> <li>(the extracting agent being alcohol or a mixture of alcohols)</li> <li>(the extracting agent being a ketone or a mixture of ketones)</li> <li>(the extracting agent being an ether or a mixture of ethers)</li> <li>(the extracting agent being an ester or a mixture of esters)</li> </ol>	31/0407	graphitic oxides, intercalated graphite, expanded graphite or graphene  NOTE  In groups C01B 31/04 - C01B 31/0492 it is desirable to add indexing codes for structural aspects or properties of graphene. The indexing codes are chosen from C01B 2204/00 - C01B 2204/32  • {Purification; Recovery or purification of graphite formed in iron making, e.g. kish graphite}  • {Intercalation}  WARNING  Group C01B 31/0415 is not complete pending a reclassification. See also group
25/463 25/464	<ol> <li>The extracting agent may be diluted with a compound or a mixture of compounds which are not solvents for phosphoric acid, e.g. a hydrocarbon</li> <li>Documents which belong to more than one subgroup of C01B 25/462 - C01B 25/466 are classified by a combination, e.g. C01B 25/462 +B4+B8</li> <li>(the extracting agent being alcohol or a mixture of alcohols)</li> <li>(the extracting agent being a ketone or a mixture of ketones)</li> <li>(the extracting agent being an ether or a mixture of ethers)</li> <li>(the extracting agent being an ester or a mixture of esters)</li> <li>(the extracting agent being a nitrogenous</li> </ol>	31/0407 31/0415	graphitic oxides, intercalated graphite, expanded graphite or graphene  NOTE  In groups C01B 31/04 - C01B 31/0492 it is desirable to add indexing codes for structural aspects or properties of graphene. The indexing codes are chosen from C01B 2204/00 - C01B 2204/32  • {Purification; Recovery or purification of graphite formed in iron making, e.g. kish graphite}  • {Intercalation}  WARNING  Group C01B 31/0415 is not complete pending a reclassification. See also group C01B 31/005
25/463 25/464 25/465	<ol> <li>The extracting agent may be diluted with a compound or a mixture of compounds which are not solvents for phosphoric acid, e.g. a hydrocarbon</li> <li>Documents which belong to more than one subgroup of C01B 25/462 - C01B 25/466 are classified by a combination, e.g. C01B 25/462 +B4+B8</li> <li>(the extracting agent being alcohol or a mixture of alcohols)</li> <li>(the extracting agent being a ketone or a mixture of ketones)</li> <li>(the extracting agent being an ether or a mixture of ethers)</li> <li>(the extracting agent being an ester or a mixture of esters)</li> <li>(the extracting agent being a nitrogenous solvent or a mixture of nitrogenous solvents</li> </ol>	31/0407 31/0415	graphitic oxides, intercalated graphite, expanded graphite or graphene  NOTE  In groups C01B 31/04 - C01B 31/0492 it is desirable to add indexing codes for structural aspects or properties of graphene. The indexing codes are chosen from C01B 2204/00 - C01B 2204/32  • {Purification; Recovery or purification of graphite formed in iron making, e.g. kish graphite}  • {Intercalation}  WARNING  Group C01B 31/0415 is not complete pending a reclassification. See also group
25/463 25/464 25/465 25/466	<ol> <li>The extracting agent may be diluted with a compound or a mixture of compounds which are not solvents for phosphoric acid, e.g. a hydrocarbon</li> <li>Documents which belong to more than one subgroup of C01B 25/462 - C01B 25/466 are classified by a combination, e.g. C01B 25/462 +B4+B8</li> <li>(the extracting agent being alcohol or a mixture of alcohols)</li> <li>(the extracting agent being a ketone or a mixture of ketones)</li> <li>(the extracting agent being an ether or a mixture of ethers)</li> <li>(the extracting agent being an ester or a mixture of esters)</li> <li>(the extracting agent being a nitrogenous solvent or a mixture of nitrogenous solvents such as amines or amides)</li> </ol>	31/0407 31/0415	graphitic oxides, intercalated graphite, expanded graphite or graphene  NOTE  In groups C01B 31/04 - C01B 31/0492 it is desirable to add indexing codes for structural aspects or properties of graphene. The indexing codes are chosen from C01B 2204/00 - C01B 2204/32  • {Purification; Recovery or purification of graphite formed in iron making, e.g. kish graphite}  • {Intercalation}  WARNING  Group C01B 31/0415 is not complete pending a reclassification. See also group C01B 31/005  • {Expanded or exfoliated graphite}
25/463 25/464 25/465	<ol> <li>The extracting agent may be diluted with a compound or a mixture of compounds which are not solvents for phosphoric acid, e.g. a hydrocarbon</li> <li>Documents which belong to more than one subgroup of C01B 25/462 - C01B 25/466 are classified by a combination, e.g. C01B 25/462 +B4+B8</li> <li>(the extracting agent being alcohol or a mixture of alcohols)</li> <li>(the extracting agent being a ketone or a mixture of ketones)</li> <li>(the extracting agent being an ether or a mixture of ethers)</li> <li>(the extracting agent being an ester or a mixture of esters)</li> <li>(the extracting agent being a nitrogenous solvent or a mixture of nitrogenous solvents such as amines or amides)</li> <li>(the extracting agent being already present</li> </ol>	31/0407 31/0415	graphitic oxides, intercalated graphite, expanded graphite or graphene  NOTE  In groups C01B 31/04 - C01B 31/0492 it is desirable to add indexing codes for structural aspects or properties of graphene. The indexing codes are chosen from C01B 2204/00 - C01B 2204/32  • {Purification; Recovery or purification of graphite formed in iron making, e.g. kish graphite}  • {Intercalation}  WARNING  Group C01B 31/0415 is not complete pending a reclassification. See also group C01B 31/005
25/463 25/464 25/465 25/466	<ol> <li>The extracting agent may be diluted with a compound or a mixture of compounds which are not solvents for phosphoric acid, e.g. a hydrocarbon</li> <li>Documents which belong to more than one subgroup of C01B 25/462 - C01B 25/466 are classified by a combination, e.g. C01B 25/462 +B4+B8</li> <li>(the extracting agent being alcohol or a mixture of alcohols)</li> <li>(the extracting agent being a ketone or a mixture of ketones)</li> <li>(the extracting agent being an ether or a mixture of ethers)</li> <li>(the extracting agent being an ester or a mixture of esters)</li> <li>(the extracting agent being a nitrogenous solvent or a mixture of nitrogenous solvents such as amines or amides)</li> <li>(the extracting agent being already present during the phosphate-containing material</li> </ol>	31/0407 31/0415	graphitic oxides, intercalated graphite, expanded graphite or graphene  NOTE  In groups C01B 31/04 - C01B 31/0492 it is desirable to add indexing codes for structural aspects or properties of graphene. The indexing codes are chosen from C01B 2204/00 - C01B 2204/32  • {Purification; Recovery or purification of graphite formed in iron making, e.g. kish graphite}  • {Intercalation}  WARNING  Group C01B 31/0415 is not complete pending a reclassification. See also group C01B 31/005  • {Expanded or exfoliated graphite}  WARNING  Groups C01B 31/043 - C01B 31/0492 are
25/463 25/464 25/465 25/466	<ol> <li>The extracting agent may be diluted with a compound or a mixture of compounds which are not solvents for phosphoric acid, e.g. a hydrocarbon</li> <li>Documents which belong to more than one subgroup of C01B 25/462 - C01B 25/466 are classified by a combination, e.g. C01B 25/462 +B4+B8</li> <li>(the extracting agent being alcohol or a mixture of alcohols)</li> <li>(the extracting agent being a ketone or a mixture of ketones)</li> <li>(the extracting agent being an ether or a mixture of ethers)</li> <li>(the extracting agent being an ester or a mixture of esters)</li> <li>(the extracting agent being a nitrogenous solvent or a mixture of nitrogenous solvents such as amines or amides)</li> <li>(the extracting agent being already present during the phosphate-containing material reaction step)</li> </ol>	31/0407 31/0415	graphitic oxides, intercalated graphite, expanded graphite or graphene  NOTE  In groups C01B 31/04 - C01B 31/0492 it is desirable to add indexing codes for structural aspects or properties of graphene. The indexing codes are chosen from C01B 2204/00 - C01B 2204/32  • {Purification; Recovery or purification of graphite formed in iron making, e.g. kish graphite}  • {Intercalation}  WARNING  Group C01B 31/0415 is not complete pending a reclassification. See also group C01B 31/005  • {Expanded or exfoliated graphite}  WARNING  Groups C01B 31/043 - C01B 31/0492 are not complete pending a reclassification. See
25/463 25/464 25/465 25/466 25/467	<ol> <li>The extracting agent may be diluted with a compound or a mixture of compounds which are not solvents for phosphoric acid, e.g. a hydrocarbon</li> <li>Documents which belong to more than one subgroup of C01B 25/462 - C01B 25/466 are classified by a combination, e.g. C01B 25/462 +B4+B8</li> <li>(the extracting agent being alcohol or a mixture of alcohols)</li> <li>(the extracting agent being a ketone or a mixture of ketones)</li> <li>(the extracting agent being an ether or a mixture of ethers)</li> <li>(the extracting agent being an ester or a mixture of esters)</li> <li>(the extracting agent being a nitrogenous solvent or a mixture of nitrogenous solvents such as amines or amides)</li> <li>(the extracting agent being already present during the phosphate-containing material reaction step)</li> <li>(the extraction being performed on the reaction</li> </ol>	31/0407 31/0415	graphitic oxides, intercalated graphite, expanded graphite or graphene  NOTE  In groups C01B 31/04 - C01B 31/0492 it is desirable to add indexing codes for structural aspects or properties of graphene. The indexing codes are chosen from C01B 2204/00 - C01B 2204/32  • {Purification; Recovery or purification of graphite formed in iron making, e.g. kish graphite}  • {Intercalation}  WARNING  Group C01B 31/0415 is not complete pending a reclassification. See also group C01B 31/005  • {Expanded or exfoliated graphite}  WARNING  Groups C01B 31/043 - C01B 31/0492 are
25/463 25/464 25/465 25/466 25/467	<ol> <li>The extracting agent may be diluted with a compound or a mixture of compounds which are not solvents for phosphoric acid, e.g. a hydrocarbon</li> <li>Documents which belong to more than one subgroup of C01B 25/462 - C01B 25/466 are classified by a combination, e.g. C01B 25/462 +B4+B8</li> <li>(the extracting agent being alcohol or a mixture of alcohols)</li> <li>(the extracting agent being a ketone or a mixture of ketones)</li> <li>(the extracting agent being an ether or a mixture of ethers)</li> <li>(the extracting agent being an ester or a mixture of esters)</li> <li>(the extracting agent being a nitrogenous solvent or a mixture of nitrogenous solvents such as amines or amides)</li> <li>(the extracting agent being already present during the phosphate-containing material reaction step)</li> <li>(the extraction being performed on the reaction slurry itself, i.e. without separating the acid</li> </ol>	31/0407 31/0415	graphitic oxides, intercalated graphite, expanded graphite or graphene  NOTE  In groups C01B 31/04 - C01B 31/0492 it is desirable to add indexing codes for structural aspects or properties of graphene. The indexing codes are chosen from C01B 2204/00 - C01B 2204/32  • {Purification; Recovery or purification of graphite formed in iron making, e.g. kish graphite}  • {Intercalation}  WARNING  Group C01B 31/0415 is not complete pending a reclassification. See also group C01B 31/005  • {Expanded or exfoliated graphite}  WARNING  Groups C01B 31/043 - C01B 31/0492 are not complete pending a reclassification. See also group C01B 31/0492 are not complete pending a reclassification. See also group C01B 31/0492 are
25/463 25/464 25/465 25/466 25/467 25/468	<ol> <li>The extracting agent may be diluted with a compound or a mixture of compounds which are not solvents for phosphoric acid, e.g. a hydrocarbon</li> <li>Documents which belong to more than one subgroup of C01B 25/462 - C01B 25/466 are classified by a combination, e.g. C01B 25/462 +B4+B8</li> <li>(the extracting agent being alcohol or a mixture of alcohols)</li> <li>(the extracting agent being a ketone or a mixture of ketones)</li> <li>(the extracting agent being an ether or a mixture of ethers)</li> <li>(the extracting agent being an ester or a mixture of esters)</li> <li>(the extracting agent being a nitrogenous solvent or a mixture of nitrogenous solvents such as amines or amides)</li> <li>(the extracting agent being already present during the phosphate-containing material reaction step)</li> <li>(the extraction being performed on the reaction slurry itself, i.e. without separating the acid (C01B 25/232 takes precedence))</li> </ol>	31/0407 31/0415	graphitic oxides, intercalated graphite, expanded graphite or graphene  NOTE  In groups C01B 31/04 - C01B 31/0492 it is desirable to add indexing codes for structural aspects or properties of graphene. The indexing codes are chosen from C01B 2204/00 - C01B 2204/32  • {Purification; Recovery or purification of graphite formed in iron making, e.g. kish graphite}  • {Intercalation}  WARNING  Group C01B 31/0415 is not complete pending a reclassification. See also group C01B 31/005  • {Expanded or exfoliated graphite}  WARNING  Groups C01B 31/043 - C01B 31/0492 are not complete pending a reclassification. See also group C01B 31/0492 are not complete pending a reclassification. See also group C01B 31/0492 are not complete pending a reclassification. See also group C01B 31/04 • {Graphitic oxides, graphitic acids or salts
25/463 25/464 25/465 25/466 25/467	<ol> <li>The extracting agent may be diluted with a compound or a mixture of compounds which are not solvents for phosphoric acid, e.g. a hydrocarbon</li> <li>Documents which belong to more than one subgroup of C01B 25/462 - C01B 25/466 are classified by a combination, e.g. C01B 25/462 +B4+B8</li> <li>(the extracting agent being alcohol or a mixture of alcohols)</li> <li>(the extracting agent being a ketone or a mixture of ketones)</li> <li>(the extracting agent being an ether or a mixture of ethers)</li> <li>(the extracting agent being an ester or a mixture of esters)</li> <li>(the extracting agent being a nitrogenous solvent or a mixture of nitrogenous solvents such as amines or amides)</li> <li>(the extracting agent being already present during the phosphate-containing material reaction step)</li> <li>(the extraction being performed on the reaction slurry itself, i.e. without separating the acid (C01B 25/232 takes precedence)</li> <li>Carbon; Compounds thereof ({C01B 6/00}),</li> </ol>	31/0407 31/0415	graphitic oxides, intercalated graphite, expanded graphite or graphene  NOTE  In groups C01B 31/04 - C01B 31/0492 it is desirable to add indexing codes for structural aspects or properties of graphene. The indexing codes are chosen from C01B 2204/00 - C01B 2204/32  • {Purification; Recovery or purification of graphite formed in iron making, e.g. kish graphite}  • {Intercalation}  WARNING  Group C01B 31/0415 is not complete pending a reclassification. See also group C01B 31/005  • {Expanded or exfoliated graphite}  WARNING  Groups C01B 31/043 - C01B 31/0492 are not complete pending a reclassification. See also group C01B 31/0492 are not complete pending a reclassification. See also group C01B 31/0492 are not complete pending a reclassification. See also group C01B 31/04  • {Graphitic oxides, graphitic acids or salts thereof}
25/463 25/464 25/465 25/466 25/467 25/468	<ol> <li>The extracting agent may be diluted with a compound or a mixture of compounds which are not solvents for phosphoric acid, e.g. a hydrocarbon</li> <li>Documents which belong to more than one subgroup of C01B 25/462 - C01B 25/466 are classified by a combination, e.g. C01B 25/462 +B4+B8</li> <li>(the extracting agent being alcohol or a mixture of alcohols)</li> <li>(the extracting agent being a ketone or a mixture of ketones)</li> <li>(the extracting agent being an ether or a mixture of ethers)</li> <li>(the extracting agent being an ester or a mixture of esters)</li> <li>(the extracting agent being a nitrogenous solvent or a mixture of nitrogenous solvents such as amines or amides)</li> <li>(the extracting agent being already present during the phosphate-containing material reaction step)</li> <li>(the extraction being performed on the reaction slurry itself, i.e. without separating the acid (C01B 25/232 takes precedence))</li> <li>Carbon; Compounds thereof ({C01B 6/00}, C01B 21/00, C01B 23/00 take precedence;</li> </ol>	31/0407	graphitic oxides, intercalated graphite, expanded graphite or graphene  NOTE  In groups C01B 31/04 - C01B 31/0492 it is desirable to add indexing codes for structural aspects or properties of graphene. The indexing codes are chosen from C01B 2204/00 - C01B 2204/32  • {Purification; Recovery or purification of graphite formed in iron making, e.g. kish graphite}  • {Intercalation}  WARNING  Group C01B 31/0415 is not complete pending a reclassification. See also group C01B 31/005  • {Expanded or exfoliated graphite}  WARNING  Groups C01B 31/043 - C01B 31/0492 are not complete pending a reclassification. See also group C01B 31/0492 (Graphitic oxides, graphitic acids or salts thereof) • {Graphene}
25/463 25/464 25/465 25/466 25/467 25/468	1. The extracting agent may be diluted with a compound or a mixture of compounds which are not solvents for phosphoric acid, e.g. a hydrocarbon  2. Documents which belong to more than one subgroup of C01B 25/462 - C01B 25/466 are classified by a combination, e.g. C01B 25/462 +B4+B8  {the extracting agent being alcohol or a mixture of alcohols}  {the extracting agent being a ketone or a mixture of ketones}  {the extracting agent being an ether or a mixture of ethers}  {the extracting agent being an ester or a mixture of esters}  {the extracting agent being a nitrogenous solvent or a mixture of nitrogenous solvents such as amines or amides}  {the extracting agent being already present during the phosphate-containing material reaction step}  {the extraction being performed on the reaction slurry itself, i.e. without separating the acid (C01B 25/232 takes precedence)}  Carbon; Compounds thereof ({C01B 6/00}, C01B 21/00, C01B 23/00 take precedence; percarbonates C01B 15/10; carbon black C09C 1/48;	31/0407	graphitic oxides, intercalated graphite, expanded graphite or graphene  NOTE  In groups C01B 31/04 - C01B 31/0492 it is desirable to add indexing codes for structural aspects or properties of graphene. The indexing codes are chosen from C01B 2204/00 - C01B 2204/32  • {Purification; Recovery or purification of graphite formed in iron making, e.g. kish graphite}  • {Intercalation}  WARNING  Group C01B 31/0415 is not complete pending a reclassification. See also group C01B 31/005  • {Expanded or exfoliated graphite}  WARNING  Groups C01B 31/043 - C01B 31/0492 are not complete pending a reclassification. See also group C01B 31/04  • {Graphitic oxides, graphitic acids or salts thereof} • {Graphene} • {Preparation}
25/463 25/464 25/465 25/466 25/467 25/468	<ol> <li>The extracting agent may be diluted with a compound or a mixture of compounds which are not solvents for phosphoric acid, e.g. a hydrocarbon</li> <li>Documents which belong to more than one subgroup of C01B 25/462 - C01B 25/466 are classified by a combination, e.g. C01B 25/462 +B4+B8</li> <li>(the extracting agent being alcohol or a mixture of alcohols)</li> <li>(the extracting agent being a ketone or a mixture of ketones)</li> <li>(the extracting agent being an ether or a mixture of ethers)</li> <li>(the extracting agent being an ester or a mixture of esters)</li> <li>(the extracting agent being a nitrogenous solvent or a mixture of nitrogenous solvents such as amines or amides)</li> <li>(the extracting agent being already present during the phosphate-containing material reaction step)</li> <li>(the extraction being performed on the reaction slurry itself, i.e. without separating the acid (C01B 25/232 takes precedence))</li> <li>Carbon; Compounds thereof ({C01B 6/00}, C01B 21/00, C01B 23/00 take precedence;</li> </ol>	31/0407	graphitic oxides, intercalated graphite, expanded graphite or graphene  NOTE  In groups C01B 31/04 - C01B 31/0492 it is desirable to add indexing codes for structural aspects or properties of graphene. The indexing codes are chosen from C01B 2204/00 - C01B 2204/32  • {Purification; Recovery or purification of graphite formed in iron making, e.g. kish graphite}  • {Intercalation}  WARNING  Group C01B 31/0415 is not complete pending a reclassification. See also group C01B 31/005  • {Expanded or exfoliated graphite}  WARNING  Groups C01B 31/043 - C01B 31/0492 are not complete pending a reclassification. See also group C01B 31/0492 (Graphitic oxides, graphitic acids or salts thereof) • {Graphene}

31/0469	• • • • {by exfoliation}		belonging to more than one of the groups
31/0409			C01B 31/303 - C01B 31/36 are classified in
	{ starting from graphitic oxide }		C01B 31/301
31/0484	{After-treatments}		<u>COID 31/301</u>
31/0492	· · · · {Purification}	31/303	• • {Simple carbides of elements not covered below}
31/06	Diamond	31/305	• • {Titanium carbides}
31/065	• • • {After-treatment, e.g. purification, irradiation}	31/306	• • {Carbides of actinides}
31/08	. Active carbon	31/308	{Carbides of alkali metals, strontium, barium
31/081	• • {from waste materials, e.g. tyres, spent sulfite		or magnesium; Mixtures thereof with calcium
	pulp liquor}		carbide}
31/082	• • {from distillation residues of coal or petroleum;	31/32	Calcium carbide
	from petroleum acid sludge}	31/34	Tungsten or molybdenum carbides
31/083	• • {After-treatment, e.g. purification (granulation	31/36	Carbides of silicon or boron
	<u>C01B 31/14</u> )}	22/00	CP C In the conf ((CO1D (100))
31/084	• • {Coating; Grafting; Microencapsulation}	33/00	Silicon; Compounds thereof ( $\{C01B 6/00\}$ ),
31/085	• • {with "molecular sieve" properties}		<u>C01B 21/00</u> , <u>C01B 23/00</u> take precedence; persilicates <u>C01B 15/14</u> ; carbides <u>C01B 31/36</u> )
31/086	• • {Preparation, reactivation or regeneration by a	22/02	
	physical process, e.g. by irradiation, by using	33/02	Silicon (forming single crystals or homogeneous  polyamytalling material with defined atmosphere
	electric current passing through a carbonaceous		polycrystalline material with defined structure
	feedstock, by using recyclable inert heating	33/021	C30B)
24/00=	bodies}	33/021	<ul> <li>Preparation (chemical coating from the vapour phase <u>C23C 16/00</u>)</li> </ul>
31/087	• • {Reactivation or regeneration (by a physical	33/023	
21/000	process <u>C01B 31/086</u> )}	33/023	by reduction of silica or {free} silica- containing material
31/088	{Apparatus (C01B 31/086 takes precedence)}	33/025	• • • • with carbon or a solid carbonaceous material,
31/089	• • {Making shaped products, e.g. fibres, spheres,	33/023	i.e. carbo-thermal process
	membranes, foam, or the like (granulation	33/027	by decomposition or reduction of gaseous or
21/10	<u>C01B 31/14</u> )}	33/027	vaporised silicon compounds other than silica
31/10	• Preparation by using gaseous activating agents		or silica-containing material
21/12	{( <u>C01B 31/086</u> , <u>C01B 31/088</u> take precedence)}	33/029	by decomposition of monosilane
31/12	• Preparation by using non-gaseous activating	33/03	by decomposition of infollostratic  by decomposition of silicon halides or
	agents {(C01B 31/086, C01B 31/088 take	33/03	halosilanes or reduction thereof with
21/125	precedence)}		hydrogen as the only reducing agent
31/125	{Preparation by impregnation with a metallic	33/031	by decomposition of silicon tetraiodide
21/14	compound}	33/031	by reduction of silicon halides or halosilanes
31/14	• • Granulation (apparatus <u>B01J 2/00</u> )	33/033	with a metal or a metallic alloy as the only
	<u>NOTE</u>		reducing agents
		33/035	by decomposition or reduction of gaseous or
	In this group, the term granulation covers the methods of preparation of active carbon using	33/033	vaporised silicon compounds in the presence
	carbonaceous precursors per se and binders,		of heated filaments of silicon, carbon or a
	e.g. pitch, and producing the granules		refractory metal, e.g. tantalum or tungsten,
	e.g. piten, and producing the granules		or in the presence of heated silicon rods
31/18	• Carbon monoxide (metal carbonyls <u>C01G</u> )		on which the formed silicon is deposited,
31/20	Carbon dioxide		a silicon rod being obtained, e.g. Siemens
31/22	Solidifying		process
31/24	<ul> <li>Methods for the preparation of carbonates or</li> </ul>	33/037	• • Purification (by zone-melting C30B 13/00)
	bicarbonates in general (percarbonates <u>C01B 15/10</u> ;	33/039	by conversion of the silicon into a compound,
	particular individual carbonates, see the relevant		optional purification of the compound, and
	groups in <u>C01B</u> - <u>C01G</u> according to the cation)		reconversion into silicon
31/26	<ul> <li>Compounds containing carbon and sulfur, e.g.</li> </ul>	33/04	<ul> <li>Hydrides of silicon</li> </ul>
	carbon disulfide, carbon oxysulfide; Thiophosgene	33/043	• • {Monosilane}
31/262	• • {Carbon disulfide}	33/046	• • {Purification}
31/265	• • • {Preparation by reacting sulfur or a sulfur	33/06	• Metal silicides (alloys <u>C22</u> )
	compound with a hydrocarbon}	33/08	. Compounds containing halogen
31/267	• • {Carbon oxysulfide}	33/10	Compounds containing silicon, fluorine, and other
31/28	. Phosgene		elements
31/30	• Carbides (alloys <u>C22</u> )	33/103	{Fluosilicic acid; Salts thereof}
31/301	• • {Oxycarbides, sulfocarbides or mixtures of	33/107	Halogenated silanes
	carbides with other bodies, e.g. graphite;	33/10705	{Tetrafluoride}
	Carbides of other non-metals, e.g. silicocarbides,	33/1071	{Tetrachloride, trichlorosilane or
	borocarbides}		silicochloroform, dichlorosilane,
	NOTE		monochlorosilane or mixtures thereof}
		33/10715	{prepared by reacting chlorine with silicon or
	Mixtures of carbides, other than those		a silicon-containing material}
	covered by group C01B 31/308, and		

33/10721	• • • • { with the preferential formation of	33/149 Coating
	tetrachloride}	33/151 by progressively adding a sol to a different
	• • • • • {from silicon}	sol, i.e. "build-up" of particles using a
33/10731	• • • • { with the preferential formation of	"heel"
22/10726	trichlorosilane}	33/152 Preparation of hydrogels
	{from silicon}	33/1525 {from or via fluosilicic acid or salts
33/10/42	• • • • {prepared by hydrochlorination of silicon or	thereof}
22/107/7	of a silicon-containing material}	33/154 by acidic treatment of aqueous silicate solutions
33/10/4/	• • • • { with the preferential formation of	
22/10752	tetrachloride}	33/1543 {using ion exchangers}
	(from silicon)	33/1546 { the first formed hydrosol being converted to a hydrogel by introduction
33/10/37	• • • • { with the preferential formation of trichlorosilane }	into an organic medium immiscible or
33/10763	• • • • • {from silicon}	only partly miscible with water}
	{Tetrabromide; Tetraiodide}	33/155 Preparation of hydroorganogels or
	{Halogenated silanes obtained by	organogels
33/10/73	disproportionation and molecular	33/157 After-treatment of gels
	rearrangement of halogenated silanes}	33/158 Purification; Drying; Dehydrating
33/10778	• • {Purification}	33/1585 {Dehydration into aerogels}
	• • · {by adsorption}	33/159 Coating or hydrophobisation
	• • • { the adsorbing material being formed in	33/16 Preparation of silica xerogels
33/10/07	situ, e.g. by partial hydrolysis}	33/163 {by hydrolysis of organosilicon compounds,
33/10794	• • • {by forming addition compounds or	e.g. ethyl orthosilicate}
33/10/71	complexes, the reactant being possibly	33/166 {by acidification of silicate in the presence of
	contained in an adsorbent}	an inert organic phase}
33/113	• Silicon oxides; Hydrates thereof {(preparing	33/18 Preparation of finely divided silica neither in
	monoxide by reduction of siliceous material	sol nor in gel form; After-treatment thereof
	<u>C01B 33/182</u> )}	(preparation of aerogels by dehydrating
33/12	Silica; Hydrates thereof, e.g. lepidoic silicic acid	gels C01B 33/158; treatment to enhance the
33/122	{Lepidoic silicic acid}	pigmenting or filling properties <u>C09C</u> )
33/124	{Preparation of adsorbing porous silica not	33/181 {by a dry process}
	in gel form and not finely divided, i.e. silicon	33/182 {by reduction of a siliceous material,
	skeletons, by acidic treatment of siliceous	e.g. with a carbonaceous reducing agent
	materials}	and subsequent oxidation of the silicon
33/126	• • • {Preparation of silica of undetermined type}	monoxide formed}
33/128	{by acidic treatment of aqueous silicate	33/183 {by oxidation or hydrolysis in the vapour
	solutions}	phase of silicon compounds such as
33/14	Colloidal silica, e.g. dispersions, gels, sols	halides, trichlorosilane, monosilane} 33/184 {by hydrolysis of tetrafluoride}
33/141	Preparation of hydrosols or aqueous	33/185 {of crystalline silica-polymorphs having
22/1/12	dispersions	molecular sieve properties, e.g. silicalites}
33/1412	• • • • {by oxidation of silicon in basic medium}	33/186 {from or via fluosilicic acid or salts thereof
33/1415	• • • • {by suspending finely divided silica in	by a wet process}
20/1415	water}	33/187 by acidic treatment of silicates
33/1417	• • • • {an aqueous dispersion being obtained}	33/193 of aqueous solutions of silicates
33/142	by acidic treatment of silicates	33/20 • Silicates (persilicates C01B 15/14; {containing
33/143	of aqueous solutions of silicates	aluminium C01B 33/26})
33/1435	{using ion exchangers}	33/22 • Magnesium silicates
33/145	Preparation of hydroorganosols, organosols	33/24 • Alkaline-earth metal silicates
22/146	or dispersions in an organic medium	33/26 • Aluminium-containing silicates, {i.e. silico-
33/146	After-treatment of sols ({preparation of	aluminates}
	hydrosols or aqueous dispersions from	33/28 {Base exchange silicates, e.g. zeolites
	hydroorganosols, organosols or dispersions in an organic medium <u>C01B 33/141</u> };	(regeneration <u>B01J 49/00</u> )}
	preparation of hydroorganosols, organosols	33/2807 {Zeolitic silicoaluminates with a
	or dispersions in an organic medium	tridimensional crystalline structure
	from hydrosols {or aqueous dispersions}	possessing molecular sieve properties;
	<u>C01B 33/145</u> )	Isomorphous compounds wherein a part
33/1465	{"Build-up" of particles using only one sol	of the aluminium ore of the silicon present
	and a "heel" consisting or not of the sol}	may be replaced by other elements such
33/148	Concentration; Drying; Dehydration;	as gallium, germanium, phosphorus;
	Stabilisation; Purification {(C01B 33/1465)	Preparation of zeolitic molecular sieves from
	takes precedence)}	molecular sieves of another type or from
33/1485	• • • • • {Stabilisation, e.g. prevention of gelling;	<pre>preformed reacting mixtures (not used, see subgroups)}</pre>
	Purification}	anogroups)}

33/2815	• • • • • { of type A (UNION CARBIDE trade name; corresponds to GRACE's types Z-12 or Z-12L)}	33/44	<ul> <li>Products obtained from layered base- exchange silicates by ion-exchange with organic compounds such as ammonium,</li> </ul>
33/2823	{from aqueous solutions of an alkali metal aluminate and an alkali metal silicate excluding any other source of		phosphonium or sulfonium compounds or by intercalation of organic compounds, e.g. organoclay material
33/283	alumina or silica} {from a reaction mixture containing	33/46	Amorphous silicates, e.g. so-called "amorphous zeolites" (crystalline zeolites <u>C01B 39/00</u> )
	at least one aluminium silicate or aluminosilicate of a clay-type, e.g. kaolin or metakaolin or its exotherm modification or allophane (containing a single clay substantially chemically modified with an acid, i.e. beyond the activation state C01B 33/2815)}	35/00	Boron; Compounds thereof (monoborane, diborane, metal borohydrides or addition complexes thereof C01B 6/00; perborates C01B 15/12; binary compounds with nitrogen C01B 21/06; {compounds of noble gases C01B 23/0005}; phosphides C01B 25/08; carbides C01B 31/36; alloys containing
33/2838	• • • • {of faujasite type, or type X or Y (UNION	25/02	boron C22)
	CARBIDE trade names; correspond	35/02	Boron; Borides
	to GRACE's types Z-14 and Z-14HS,	35/023	{Boron}
	respectively)}	35/026	<ul> <li>{Higher boron hydrides, i.e. containing at least three boron atoms}</li> </ul>
33/2846	$\ldots$ { of type X}	35/04	Metal borides
33/2853	$\dots$ { of type Y }	35/04	
33/2861	• • • • {of mordenite type, e.g. ptilolite or		Boron halogen compounds  (Helidae)
22/20/20	dachiardite}	35/061	{Halides}
33/2869	(of other types characterised by an X-ray	35/063	<ul><li>. {Tetrafluoboric acid; Salts thereof}</li><li> {Tetrafluoboric acid}</li></ul>
22/2076	spectrum and a definite composition}	35/065 35/066	{Alkali metal tetrafluoborates}
33/2876	{from a reacting mixture containing an amine or an organic cation, e.g. a	35/068	{Arkan metal terrandoborates}     {Halogenated hydrides}
	quaternary onium cation-ammonium,	35/008	<ul> <li>• {Halogenated hydrides}</li> <li>• Compounds containing boron and nitrogen,</li> </ul>
22/2004	phosphonium, stibonium}		phosphorus, oxygen, sulfur, selenium or tellurium
33/2884	{ the aluminium or the silicon in the network being partly replaced }	35/10	Compounds containing boron and oxygen
33/2892	• • • • {containing an element or a compound	25/1000	(C01B 35/06 takes precedence)
33/2092	occluded in the pores of the network, e.g.	35/1009	• • {having molecular-sieve properties}
	an oxide already present in the starting	35/1018	<ul> <li>. (Carbonyl compounds derived from boron hydrides)</li> </ul>
	reaction mixture}	35/1027	· · · {Oxides}
33/32	• Alkali metal silicates ({C01B 33/24},	35/1027	
	C01B 33/26 take precedence)	35/1036	{Boric anhydride}
33/325	• • • { After-treatment, e.g. purification or	35/1043	<ul><li> {Oxyacids}</li><li> {Orthoboric acid}</li></ul>
	stabilsation of solutions, granulation;	35/1034	• • • {Orthoporte acid} • • • • {Preparation from boron ores or borates
	Dissolution; Obtaining solid silicate, e.g. from	33/1003	using acids or salts}
	a solution by spray-drying, flashing off water	35/1072	{by means of ammonia-carbon dioxide}
	or adding a coagulant}	35/1072	{Preparation by working up other natural
	NOTE		sources, e.g. seawater}
	In this group, obtaining solid silicate, e.g.	35/109	{Purification; Separation; Concentration}
	as a hydrate of a crystalline silicate, from	35/12	• • Borates {( <u>C01B 35/1063</u> takes precedence)}
	a solution or a hydrate melt by heating or	35/121	• • • {of alkali metal}
	cooling with or without seeding, is not	35/122	• • • • {Sodium tetraborates; Hydrates thereof,
	considered as after-treatment, but classified in group C01B 33/32	35/123	e.g. borax} {Preparation from boron ores or other
33/36	having base-exchange properties but not having	25/124	borates}
	molecular sieve properties (regeneration thereof B01J 49/00)	35/124	• • • • • {Preparation by working up natural brines, e.g. seawater}
33/38	Layered base-exchange silicates, e.g. clays, micas or alkali metal silicates of kenyaite	35/125	{Purification; Concentration; Dehydration; Stabilisation; Other after- treatment}
	or magadiite type {(activation of naturally occurring clays <u>B01J 20/12</u> ; pillared layered	35/126	• • • { of alkaline-earth metals, beryllium, aluminium or magnesium}
22/42	base-exchange silicates <u>B01J 29/049</u> )}	35/127	• • • • {of heavy metals}
33/40	Clays	35/128	{containing plural metal or metal and
33/405	{not containing aluminium}		ammonium}
33/42	Micas; {Interstratified clay-mica products (delaminated mica or vermiculite platelets	35/14	Compounds containing boron and nitrogen, phosphorus, sulfur, selenium or tellurium
	obtained by a process involving cation-	35/143	• • • {Phosphates}
33/425	exchange C04B 14/208)} {not containing aluminium}		, ,
33/423	Inot containing aruminum;		

35/146	• • • {Compounds containing boron and nitrogen, e.g. borazoles (ammonium tetrafluoborates <u>C01B 35/063</u> ; ammonium borates <u>C01B 35/12</u> )}	39/023	• • {Preparation of physical mixtures or intergrowth products of zeolites chosen from group C01B 39/04 or two or more of groups C01B 39/14 - C01B 39/48}
	001dt65 <u>0012 55/12</u> /)	39/026	• • {After-treatment}
Compounds	characterised primarily by their physical or	39/04	using at least one organic template directing
	Operties, rather than by their chemical constitution	2370.	agent, e.g. an ionic quaternary ammonium compound or an aminated compound
37/00	Compounds having molecular sieve properties but not having base-exchange properties	39/06	Preparation of isomorphous zeolites characterised by measures to replace the aluminium or silicon
	NOTE		atoms in the lattice framework by atoms of other elements, {i.e. by direct or secondary synthesis}
	Compounds classified in main group C01B 37/00 are also classified in other groups of class C01 according to their composition	39/065	• • • {Galloaluminosilicates; Group IVB-metalloaluminosilicates; Ferroaluminosilicates}
27/002	(Motellanhoenhotes not containing aluminium a g	39/08	• • • the aluminium atoms being wholly replaced
37/002	• {Metallophosphates not containing aluminium, e.g.	39/082	{Gallosilicates}
27/005	gallophosphates or silicogallophosphates}	39/085	• • • • {Group IVB- metallosilicates}
37/005	• {Silicates, i.e. so-called metallosilicalites or	39/087	• • • {Ferrosilicates}
27/007	metallozeosilites}	39/10	• • • the replacing atoms being {at least} phosphorus
37/007	• {Borosilicates}		atoms
37/02	Crystalline silica-polymorphs, e.g. silicalites {dealuminated aluminosilicate zeolites}	39/12	<ul><li>the replacing atoms being {at least} boron atoms</li></ul>
37/04	<ul> <li>Aluminophosphates (APO compounds)</li> </ul>	39/14	Type A
37/06	Aluminophosphates containing other elements, e.g. metals, boron	39/145	<ul><li>• (using at least one organic template directing agent)</li></ul>
37/065	• • {the other elements being metals only}	39/16	from aqueous solutions of an alkali metal
37/08	• Silicoaluminophosphates (SAPO compounds), {e.g. CoSAPO}		aluminate and an alkali metal silicate excluding any other source of alumina or silica but seeds
39/00	Compounds having molecular sieve and base-	20/10	$\{(\underline{\text{C01B } 39/145} \text{ takes precedence})\}$
	exchange properties, e.g. crystalline zeolites; Their preparation; After-treatment, e.g. ion-exchange or dealumination (treatment to modify the sorption properties, e.g. shaping using a binder, <u>B01J 20/10</u> ; treatment to modify the catalytic properties, e.g.	39/18	• • • from a reaction mixture containing at least one aluminium silicate or aluminosilicate of a clay type, e.g. kaolin or metakaolin or its exotherm modification or allophane {(C01B 39/145 takes precedence)}
	combination of treatments to make the zeolites	39/20	• • Faujasite type, e.g. type X or Y
	appropriate to their use as a catalyst, <u>B01J 29/04</u> ; treatment to improve the ion-exchange properties	39/205	• • • {using at least one organic template directing agent; Hexagonal faujasite; Intergrowth
	<u>B01J 39/14</u> ; regeneration or reactivation of ion-	20/22	products of cubic and hexagonal faujasite}
	exchange properties <u>B01J 49/00</u> ; preparation of	39/22	Type X { $(C01B 39/205 \text{ takes precedence})$ }
	stabilised suspensions used in detergents <u>C11D 3/12</u> )	39/24	Type Y $\{(\underline{\text{C01B } 39/205} \text{ takes precedence})\}$
	NOTES	39/26	• Mordenite type {(C01B 39/023, C01B 39/026, C01B 39/06 take precedence)}
	1. In this group, the following term is used with the meaning indicated:	39/265	• • • {using at least one organic template directing agent}
	<ul><li> "zeolites" means:</li><li> i. crystalline aluminosilicates with base-</li></ul>	39/28	• Phillipsite or harmotome type {( <u>C01B 39/023</u> , <u>C01B 39/026</u> , <u>C01B 39/06</u> take precedence)}
	exchange and molecular sieve properties,	39/30	• • Erionite or offretite type, e.g. zeolite T
	having three dimensional, microporous lattice framework structure of tetrahedral	39/305	<ul> <li> {using at least one organic template directing agent}</li> </ul>
	oxide units;	39/32	Type L
	ii. compounds isomorphous to those of the	39/34	Type ZSM-4
	former category, wherein the aluminium or silicon atoms in the framework are partly or wholly replaced by atoms of other elements,	39/36	<ul> <li>Pentasil type, e.g. types ZSM-5, ZSM-8 or ZSM-11</li> </ul>
	e.g. by gallium, germanium, phosphorus or boron.	39/365	• • • {Type ZSM-8; Type ZSM-11; ZSM 5/11 intermediate}
		39/38	Type ZSM-5
	2. Compounds classified in main group C01B 39/00 are also classified in other groups of class C01	39/40	• • • using at least one organic template directing agent
	according to their composition	39/42	Type ZSM-12
39/02	<ul> <li>Crystalline aluminosilicate zeolites; Isomorphous compound thereof; Direct preparation thereof;</li> </ul>	39/44	<ul> <li>Ferrierite type, e.g. types ZSM-21, ZSM-35 or ZSM-38</li> </ul>
	Preparation thereof starting from a reaction mixture containing a crystalline zeolite of another type, or from preformed reactants; After-treatment thereof	39/445	• • • {using at least one organic template directing agent}

Integrated processes for the production of

B01J 2208/00 - B01J 2219/00) 2203/02 • Processes for making hydrogen or synthesis gas

2203/0211 . . . containing a non-catalytic reforming step 2203/0216 . . . containing a non-catalytic steam reforming

2203/0222 . . . containing a non-catalytic carbon dioxide

2203/0233 . . . the reforming step being a steam reforming

2203/0238 . . . the reforming step being a carbon dioxide

reforming step

2203/0244 . . . the reforming step being an autothermal

reforming step

2203/0227 . . . containing a catalytic reforming step

2203/0205 . . containing a reforming step

step

hydrogen or synthesis gas (reactors or details thereof

reforming step, e.g. secondary reforming

2203/00

39/46	<ul> <li>Other types characterised by their X-ray</li> </ul>
	diffraction pattern and their defined composition
	{( <u>C01B 39/023</u> , <u>C01B 39/026</u> , <u>C01B 39/06</u> take
	precedence)}
39/48	using at least one organic template directing
	agent
39/50	. Zeolites wherein inorganic bases or salts occlude
	channels in the lattice framework e.g. sodalite

channels in the lattice framework, e.g. sodalite, cancrinite, nosean, hauynite {(ultramarine C09C 1/32)}

39/52 Sodalites

2202/36 . . Diameter

39/54

. Phosphates, e.g. APO or SAPO compounds

## **NOTE**

Phosphates having either a poorly defined or a weak base-exchange capacity such as MAPO's, SAPO's or BAPO's are classified in C01B 37/00

		processes
		2203/025 containing a partial oxidation step
		2203/0255 containing a non-catalytic partial oxidation step
2201/00	Preparation of ozone by electrical discharge	2203/0261 containing a catalytic partial oxidation step
2201/10	Dischargers used for production of ozone	[CPO]
2201/12	Plate-type dischargers	2203/0266 containing a decomposition step
2201/14	Concentric/tubular dischargers	2203/0272 containing a non-catalytic decomposition step
2201/20	Electrodes used for obtaining electrical discharge	2203/0277 containing a catalytic decomposition step
2201/22	• Constructional details of the electrodes	2203/0283 containing a CO-shift step, i.e. a water gas shift
2201/24	• Composition of the electrodes	step
2201/30	Dielectrics used in the electrical dischargers	2203/0288 containing two CO-shift steps
2201/32	Constructional details of the dielectrics	2203/0294 containing three or more CO-shift steps
2201/34	Composition of the dielectrics	2203/04 . containing a purification step for the hydrogen or
2201/40	<ul> <li>using several dischargers in series</li> </ul>	the synthesis gas
2201/50	Part of the product being recycled	2203/0405 Purification by membrane separation
2201/60	Feed streams for electrical dischargers	2203/041 In-situ membrane purification during hydrogen
2201/62	Air	production
2201/64	Oxygen	2203/0415 . Purification by absorption in liquids
2201/66	Pretreatment of the feed	2203/042 • Purification by adsorption on solids
2201/70	Cooling of the discharger; Means for making	2203/0425 In-situ adsorption process during hydrogen
	cooling unnecessary	production
2201/72	by air	2203/043 Regenerative adsorption process in two or
2201/74	by liquid	more beds, one for adsorption, the other for
2201/76	Water	regeneration
2201/80	Additional processes occurring alongside the	2203/0435 Catalytic purification
	electrical discharges, e.g. catalytic processes	2203/044 Selective oxidation of carbon monoxide
2201/82	Treatment with ultraviolet light	2203/0445 Selective methanation
2201/84	Treatment with magnetic fields	2203/045 Purification by catalytic desulfurisation
2201/90	• Control of the process	2203/0455 . Purification by non-catalytic desulfurisation
2202/00	Characteristics of coulon monotoles	2203/046 • Purification by cryogenic separation
2202/00	Structure or properties of carbon nanotubes	2203/0465 Composition of the impurity
2202/02	Single-walled nanotubes	2203/047 the impurety being carbon monoxide
2202/04	Nanotubes with a specific amount of walls	2203/0475 the impurety being carbon dioxide
2202/06	Multi-walled nanotubes	2203/048 the impurety being an organic compound
2202/08	Aligned nanotubes	2203/0485 the impurety being a sulfur compound
2202/10	• Filled nanotubes	2203/049 the impurety being carbon
2202/20	Nanotubes characterized by their properties	2203/0495 the impurety being water
2202/22	Electronic properties	• Integration with other chemical processes
2202/24	Thermal properties	2203/061 Methanol production
2202/26	Mechanical properties	2203/062 Hydrocarbon production, e.g. Fischer-Tropsch
2202/28	Solid content in solvents	process
2202/30	Purity	2203/063 . Refinery processes
2202/32	Specific surface area	2203/065 using hydrotreating, e.g. hydrogenation,
2202/34	Length	hydrodesulfurisation

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2203/066 . . with fuel cells

2203/067 the reforming process taking place in the fuel	2203/1264	Catalytic pre-treatment of the feed
cell	2203/127	Catalytic desulfurisation
2203/068 Ammonia synthesis	2203/1276	Mixing of different feed components
2203/08 • Methods of heating or cooling	2203/1282	using static mixers
2203/0805 Methods of heating the process for making hydrogen or synthesis gas	2203/1288	Evaporation of one or more of the different feed
2203/0811 by combustion of fuel	2202/1204	components
•	2203/1294	• Evaporation by heat exchange with hot process
2203/0816 Heating by flames	2202/14	stream
2203/0822 the fuel containing hydrogen	2203/14	Details of the flowsheet
2203/0827 at least part of the fuel being a recycle stream	2203/141	. At least two reforming, decomposition or partial
2203/0833 Heating by indirect heat exchange with hot	2202/142	oxidation steps in parallel
fluids, other than combustion gases, product gases or non-combustive exothermic reaction	2203/142	At least two reforming, decomposition or partial oxidation steps in series
product gases	2203/143	Three or more reforming, decomposition or
2203/0838 by heat exchange with exothermic reactions,	2200/110	partial oxidation steps in series
other than by combustion of fuel	2203/145	At least two purification steps in parallel
2203/0844 the non-combustive exothermic reaction	2203/146	At least two purification steps in series
being another reforming reaction as defined	2203/147	Three or more purification steps in series
in groups <u>C01B 2203/02</u> - <u>C01B 2203/0294</u>	2203/148	. involving a recycle stream to the feed of the
2203/085 by electric heating	2203/110	process for making hydrogen or synthesis gas
2203/0855 by electromagnetic heating	2203/16	• Controlling the process
2203/0861 by plasma	2203/1604	Starting up the process
2203/0866 by combination of different heating methods	2203/1609	Shutting down the process
2203/0872 Methods of cooling	2203/1614	Controlling the temperature
2203/0877 by direct injection of fluid	2203/1619	Measuring the temperature
2203/0883 by indirect heat exchange	2203/1613	Adjusting the temperature
2203/0888 by evaporation of a fluid	2203/1628	Controlling the pressure
2203/0894 Generation of steam	2203/1628	Measuring the pressure
2203/10 • Catalysts for performing the hydrogen forming	2203/1638	Adjusting the pressure
reactions	2203/1638	Controlling the product
2203/1005 Arrangement or shape of catalyst		
2203/1011 • • • Packed bed of catalytic structures, e.g.	2203/1647	Controlling the amount of the product
particles, packing elements	2203/1652	Measuring the amount of product
2203/1017 characterised by the form of the structure	2203/1657	the product being hydrogen
2203/1023 Catalysts in the form of a monolith or	2203/1661	the product being carbon monoxide
honeycomb	2203/1666	the product being carbon dioxide
2203/1029 Catalysts in the form of a foam	2203/1671	Controlling the composition of the product
2203/1035 Catalyst coated on equipment surfaces, e.g.	2203/1676	Measuring the composition of the product
reactor walls	2203/168	Adjusting the composition of the product
2203/1041 Composition of the catalyst	2203/1685	Control based on demand of downstream process
2203/1047 Group VIII metal catalysts	2203/169	Controlling the feed
2203/1052 Nickel or cobalt catalysts	2203/1695	Adjusting the feed of the combustion
2203/1058 Nickel catalysts	2203/80	Aspect of integrated processes for the production
2203/1064 Platinum group metal catalysts		of hydrogen or synthesis gas not covered by groups
2203/107 Platinum catalysts	2202/92	C01B 2203/02 - C01B 2203/1695  • Several process steps of
2203/1076 Copper or zinc-based catalysts	2203/82	C01B 2203/02 - C01B 2203/08 integrated into a
2203/1082 Composition of support materials		single apparatus
2203/1088 Non-supported catalysts	2203/84	Energy production
2203/1094 Promotors or activators	2203/86	Carbon dioxide sequestration
2203/12 • Feeding the process for making hydrogen or	2203/00	· · Carbon dioxide sequestration
synthesis gas	2204/00	Structure or properties of graphene
2203/1205 Composition of the feed	2204/02	Single layer graphene
2203/1211 Organic compounds or organic mixtures used	2204/04	Specific amount of layers or specific thickness
in the process for making hydrogen or synthesis	2204/06	Graphene nanoribbons
gas	2204/065	characterized by their width or by their aspect
2203/1217 Alcohols		ratio
2203/1223 Methanol	2204/20	Graphene characterized by its properties
2203/1229 Ethanol	2204/22	Electronic properties
2203/1235 Hydrocarbons	2204/24	Thermal properties
2203/1241 Natural gas or methane	2204/26	Mechanical properties
2203/1247 Higher hydrocarbons	2204/28	Solid content in solvents
2203/1252 Cyclic or aromatic hydrocarbons	2204/30	Purity
2203/1258 • Pre-treatment of the feed		

2204/32	Size or surface area
2210/00	Purification or separation of specific gases
2210/0001	Separation or purification processing
2210/0003	Chemical processing
2210/0004	by oxidation
2210/0006	by reduction
2210/0007	by complexation
2210/0009	Physical processing
2210/001	by making use of membranes
2210/0012	characterised by the membrane
2210/0014	by adsorption in solids
2210/0015	characterised by the adsorbent
2210/0017	Carbon-based materials
2210/0018	Zeolites
2210/002	Other molecular sieve materials
2210/0021	Temperature swing adsorption
2210/0023	in getters
2210/0025	by absorption in liquids
2210/0026	Isotopes of the specific gas
2210/0028	<ul> <li>Separation of the specific gas from gas mixtures</li> </ul>
	containing a minor amount of this specific gas
2210/0029	• Obtaining noble gases
2210/0031	Helium
2210/0032	Neon
2210/0034	Argon
2210/0035	Krypton
2210/0037	Xenon
2210/0039	Radon
2210/004	Separation of a mixture of noble gases
2210/0042	Making ultrapure specific gas
2210/0043	Impurity removed
2210/0045	. Oxygen
2210/0046	. Nitrogen
2210/0048	Air
2210/005	Carbon monoxide
2210/0051	Carbon dioxide
2210/0053	Hydrogen
2210/0054	Hydrogen halides
2210/0056	Hydrogen fluoride
2210/0057	Hydrogen chloride
2210/0059	Hydrogen bromide
2210/006	Hydrogen iodide
2210/0062	Water
2210/0064	Hydrogen sulfide
2210/0065	. Ammonia
2210/0067	Hydrogen cyanide
2210/0068	Organic compounds
2210/007	Hydrocarbons
2210/0071	Sulfur oxides
2210/0073	Sulfur halides
2210/0075	Nitrogen oxides
2210/0076	Nitrogen halides
2210/0078	Noble gases
2210/0079	Helium
2210/0077	Neon
2210/0081	Argon
2210/0082	Krypton
2210/0084	Xenon
2210/0083	Radon
2210/0087	. Peroxides
2210/0089	Hydrogen peroxide
2210/009	• • • Trydrogen peroxide

2210/0092 . . Ozone
2210/0093 . . Metals or metal compounds
2210/0095 . . . Metals
2210/0096 . . . Metal hydrides

2210/0098 . . Other impurities